

STIC-Biotech/ChemLib

60001

From: Ramirez, Delia
Sent: Friday, February 08, 2002 7:08 PM
To: STIC-Biotech/ChemLib
Subject: case 09/606129

Hi,

I would like to request the following searches (09/606129 Maines et al.)

1. a standard search of seq id 1, 3, 18, 19, 34, 35 in the protein databases (commercial and interference)
2. an oligo search of seq id 18, 19, 34, 35 in the protein databases (commercial)

Thank you,

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02/08/02

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Searcher: _____
Phone: _____
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Date Picked Up: 2/12/02
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Clerical: _____
Online time: _____

TYPE OF SEARCH:
NA Sequences: _____
AA Sequences: 10
Structures: _____
Bibliographic: _____
Litigation: _____
Full text: _____
Patent Family: _____
Other: _____

VENDOR/COST(where applic.)
STN: _____
DIALOG: _____
Questel/Orbit: _____
DRLink: _____
Lexis/Nexis: _____
Sequence Sys.: 02
WWW/Internet: _____
Other (specify): _____

1

Pending Nucleic Acid and/or Pending Amino Acid database searches now generate two sets of results. These databases were split into two parts to reduce the time needed to update the databases daily. The split freed up more machine time for processing searches.

Searches run against the Nucleic Acid Pending database produce two sets of results, with the extensions, **.rnpm** and **.rnpn**

Searches run against the Amino Acid Pending database produce two sets of results, with the extensions, **.rapm** and **.rapn**

The Pending database search results should not be left in the case because they contain data that is confidential.

GenCore version 4.5
Copyright (c) 1993 - 2000 CompuGen Ltd.

OM protein - protein search, using sw model

Run on: February 12, 2002, 12:03:24 ; Search time 94.82 Seconds
(without alignments)
10.798 Million cell updates/sec

Title: US-09-606-129A-18

Perfect score: 41

Sequence: 1 KKRIMHC 7

Scoring table: BLOSUM62
Gapop 10.0 , Gapext 0.5

Searched: 473505 segs, 146272329 residues

Total number of hits satisfying chosen parameters: 473505

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database :

SPTREMBL_17:*
1: sp_archaea:*
2: sp_bacteria:*
3: sp_fungi:*
4: sp_human:*
5: sp_invertebrate:*
6: sp_mammal:*
7: sp_mhc:*
8: sp_organelle:*
9: sp_phage:*
10: sp_plant:*
11: sp_rodent:*
12: sp_virus:*
13: sp_vertebrate:*
14: sp_unclassified:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	41	100.0	599	10 Q9FHV3	Q9fHV3 arabidopsis
2	41	100.0	633	10 Q49511	Q49511 arabidopsis
3	38	92.7	295	11 Q9CY64	Q9cy64 mus musculus
4	38	92.7	296	4 Q9BRW8	Q9brw8 homo sapien
5	38	92.7	303	11 Q9DD21	Q9dd21 mus musculus
6	37	90.2	106	5 Q9XZM5	Q9xxm5 leishmania
7	36	87.8	512	10 Q9LML0	Q9lml0 arabidopsis
8	35	85.4	508	11 Q9P6S2	Q9d6s2 mus musculus
9	34	82.9	279	11 Q9CXV3	Q9cxv3 mus musculus
10	34	82.9	401	10 Q9M149	Q9m149 arabidopsis
11	34	82.9	431	10 Q04613	Q04613 arabidopsis
12	33	80.5	189	2 Q03949	Q03949 anabaena sp
13	33	80.5	267	2 Q9AKS0	Q9aks0 pseudomonas
14	33	80.5	495	10 Q9ST63	Q9st63 solanum tub
15	33	80.5	508	10 Q80874	Q80874 arabidopsis
16	32	78.0	102	10 Q9S738	Q9s738 lycopersico
17	32	78.0	104	12 Q9Q553	Q9q553 human immun
18	32	78.0	108	2 P75909	P75909 escherichia
19	32	78.0	111	4 Q9H007	Q9h007 homo sapien

ALIGNMENTS

RESULT 1
Q9FHV3
ID Q9FHV3 PRELIMINARY; PRT; 599 AA.
AC Q9FHV3;
DT 01-MAR-2001 (TRENBLrel. 16, Created)
DT 01-MAR-2001 (TRENBLrel. 16, Last sequence update)
DE 01-MAR-2001 (TRENBLrel. 16, Last annotation update)
DE MUTATOR-LIKE TRANSPOSASE-LIKE PROTEIN.
OS Arabidopsis thaliana (Mouse-ear cress).
OC Eukaryota; Viridiplantae; Streptophyta;
OC Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots; Rosidae;
OC eurosids II; Brassicales; Brassicaceae; Arabidopsi.
OX NCBI_taxid=3702;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN-COLUMBIA;
RX MEDLINE=99397451; PubMed=10470850;
RA Kaneko T., Katoh T., Sato S., Nakamura Y., Asamizu E., Kotani H.,
RA Miyajima N., Tabata S.;
RT "Structural analysis of Arabidopsis thaliana chromosome 5. IX.
RT Sequence features of the regions of 1,011,550 bp covered by seventeen
RT pl and TAC clones.";
RL DNA Res. 6:183-195(1999).
DR EMBL; AB017068; BAB11366.1;
SQ SEQUENCE 599 AA; 69407 MW; 36BAEE2F2A82D717 CRC64;

Query Match 100.0%; Score 41; DB 10; Length 599;
Best Local Similarity 100.0%; Pred. No. 0.73;
Matches 7; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 KKRIMHC 7

Db 563 KKRIMHC 569

RESULT 2

Q49511 PRELIMINARY; PRT; 633 AA.
ID Q49511
AC Q49511;
DT 01-JUN-1998 (TRENBLrel. 06, Created)
DT 01-JUN-1998 (TRENBLrel. 06, Last sequence update)

32 78.0 142 4 Q9BTS7
32 78.0 233 4 Q9UIJ2
32 78.0 235 4 Q9P0U6
32 78.0 279 4 Q43658
32 78.0 279 4 Q9UK52
32 78.0 279 4 Q9HD98
32 78.0 283 4 Q9HD99
32 78.0 294 4 Q00532
32 78.0 338 3 Q9CIL3
32 78.0 377 2 Q9KSC1
32 78.0 422 4 Q9UIJ4
32 78.0 422 4 Q9UK53
32 78.0 455 5 Q9VJ28
32 78.0 497 10 Q9C8V1
32 78.0 516 5 Q21592
32 78.0 571 10 Q24022
32 78.0 601 4 Q9BRB5
32 78.0 601 4 Q9U6W5
31 75.6 109 5 Q9VZM4
31 75.6 137 4 Q9NSR1
31 75.6 152 5 Q44797
31 75.6 279 2 Q9AJ03
31 75.6 321 10 Q9LZD9
31 75.6 398 5 Q9VT97
31 75.6 446 2 Q9CKP7
31 75.6 476 4 Q9UFK0

Q9bts7 homo sapien
Q9uij2 homo sapien
Q9p0u6 homo sapien
Q43658 homo sapien
Q9uk52 homo sapien
Q9hd98 homo sapien
Q9hd99 homo sapien
Q00532 homo sapien
Q9cil3 neosporea
Q9kscl vibrio chol
Q9uij4 homo sapien
Q9uk53 homo sapien
Q9vj28 drosophila
Q9c8v1 arabidopsis
Q21592 caenorhabdi
Q24022 lycopersico
Q9brb5 homo sapien
Q9u6w5 caenorhabdi
Q9vzm4 drosophila
Q9nsr1 homo sapien
Q44797 caenorhabdi
Q44797 caenorhabdi
Q9aj03 staphylococ
Q9lzd9 arabidopsis
Q9vt97 drosophila
Q9ckp7 pasteurella
Q9ufk0 homo sapien

RA Sakai K., Okido T., Furuno M., Aono H., Baldarelli R., Barsh G.,
 RA Blake J., Boffelli D., Bojunga N., Carninci P., de Bonaldo M.F.,
 RA Brownstein M.J., Bult C., Fletcher C., Fujita M., Gariboldi M.,
 RA Gustincich S., Hill D., Hofmann M., Hume D.A., Kamiya M., Lee N.H.,
 RA Lyons P., Marchionni L., Mashima J., Mazzarelli J., Mombaerts P.,
 RA Nordone P., Ring B., Ringwald M., Rodriguez I., Sakamoto N.,
 RA Sasaki H., Sato K., Schoenbach C., Seya T., Shibata Y., Storch K.-F.,
 RA Suzuki H., Toyooka K., Wang K.H., Weitz C., Whittaker C., Wilming L.,
 RA Wainshaw-Boris A., Yoshida K., Hasegawa Y., Kawaji H., Kohsaki S.,
 RA Hayashizaki Y.,
 RT "Functional annotation of a full-length mouse cDNA collection.";
 RL Nature 409:685-690(2001).
 DR EMBL; AK002231; BAB21950.1;
 DR MGD; MGI:1915580; 0610006A11rik.
 DR InterPro; IPR000683; GFO_IDH_Moca.
 DR Pfam; PF01408; GFO_IDH_Moca; 1.
 SQ SEQUENCE 303 AA; 34491 MW; 52DBA3B02EB956EB CRC64;

Query Match 92.7%; Score 38; DB 11; Length 303;
 Best Local Similarity 85.7%; Pred. No. 1.8;
 Matches 6; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

QY 1 KKRIMHC 7
 |||||
 Db 274 KKRILHC 280

RESULT 6
 Q9XZM5 PRELIMINARY; PRT; 106 AA.
 AC Q9XZM5;
 DT 01-NOV-1999 (TrEMBLrel. 12, Created)
 DT 01-MAY-2000 (TrEMBLrel. 13, Last sequence update)
 DT 01-JUN-2001 (TrEMBLrel. 17, Last annotation update)
 DE RIBOSOMAL PROTEIN L44.
 GN RPL44.
 OS Leishmania amazonensis.
 OC Eukaryota; Euglenozoa; Kinetoplastida; Trypanosomatidae; Leishmania.
 ON NCBI_TaxID=5659;
 RP SEQUENCE FROM N.A.
 RC STRAIN=LV78;
 RT "Cloning and characterization of Leishmania ribosomal protein L44.";
 RL Submitted (MAR-2000) to the EMBL/GenBank/DBJ databases.
 CC -!- SIMILARITY: TO THE L44E FAMILY OF RIBOSOMAL PROTEINS.
 DR EMBL; AF148853; AAD31928.2;
 DR InterPro; IPR000552; Ribosomal_L44E.
 DR Pfam; PF00935; Ribosomal_L44; 1.
 DR ProDom; PD002841; Ribosomal_L44E; 1.
 DR PROSITE; PS01172; RIBOSOMAL_L44E; 1.
 KW Ribosomal protein.
 SQ SEQUENCE 106 AA; 12283 MW; F30A3AB2047B0334 CRC64;

Query Match 90.2%; Score 37; DB 5; Length 106;
 Best Local Similarity 71.4%; Pred. No. 1.2;
 Matches 5; Conservative 2; Mismatches 0; Indels 0; Gaps 0;

QY 1 KKRIMHC 7
 |||||
 Db 6 KKKVMHC 12

RESULT 7
 Q9LML0 PRELIMINARY; PRT; 512 AA.
 AC Q9LML0;
 DT 01-OCT-2000 (TrEMBLrel. 15, Created)
 DT 01-OCT-2000 (TrEMBLrel. 15, Last sequence update)
 DT 01-JUN-2001 (TrEMBLrel. 17, Last annotation update)
 DE F10K1.10 PROTEIN.

GN F10K1.10.
 OS Arabidopsis thaliana (Mouse-ear cress).
 OC Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
 OC Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots; Rosidae;
 OC eurosids II; Brassicales; Brassicaceae; Arabidopsids.
 OX NCBI_TaxID=3702;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN=CV. COLUMBIA;
 RA Liu S.X., Chan A., Yu G., Etgu P., Lee J.M., Lenz C., Pham P.,
 RA Sakano H., Toriumi M., Chung M., Goldsmith A., Liu A., Smith A.,
 RA Vaysberg M., Altafi H., Brooks S., Buehler E., Chao Q., Conn L.,
 RA Conway A., Hansen N., Johnson-Hopson C., Kian S., Kim C., Lam B.,
 RA Miranda M., Nguyen M., Palm C.J., Shinn P., Southwick A., Davis R.W.,
 RA Ecker J.R., Federspiel N.A., Theologis A.;
 RT "The sequence of BAC F10K1 from Arabidopsis thaliana chromosome 1.";
 RL Submitted (APR-2000) to the EMBL/GenBank/DBJ databases.
 RN [2]
 RP SEQUENCE FROM N.A.
 RC STRAIN=CV. COLUMBIA;
 RA Theologis A.;
 RL Submitted (APR-2000) to the EMBL/GenBank/DBJ databases.
 RN [3]
 RP SEQUENCE FROM N.A.
 RC STRAIN=CV. COLUMBIA;
 RA Theologis A.;
 RL Submitted (JUN-2000) to the EMBL/GenBank/DBJ databases.
 RN [4]
 RP SEQUENCE FROM N.A.
 RC STRAIN=CV. COLUMBIA;
 RA Theologis A.;
 RL Submitted (JUL-2000) to the EMBL/GenBank/DBJ databases.
 CC -!- COFACTOR: FAD (BY SIMILARITY).
 CC -!- SIMILARITY: TO PYRIDINE NUCLEOTIDE-DISULPHIDE OXIDOREDUCTASES
 CC CLASS=1.
 DR EMBL; AC067971; AAF92202.1;
 DR InterPro; IPR001327; FAD_pyr_redox.
 DR Pfam; PF00070; pyr_redox; 1.
 KW FAD; Flavoprotein; Oxidoreductase; Redox-active center.
 SQ SEQUENCE 512 AA; 56857 MW; 1F63AFA9A1A2C13B CRC64;

Query Match 87.8%; Score 36; DB 10; Length 512;
 Best Local Similarity 71.4%; Pred. No. 7.5;
 Matches 5; Conservative 2; Mismatches 0; Indels 0; Gaps 0;

QY 1 KKRIMHC 7
 |||||
 Db 236 KKRILHC 242

RESULT 8
 Q9D6S2 PRELIMINARY; PRT; 508 AA.
 AC Q9D6S2;
 DT 01-JUN-2001 (TrEMBLrel. 17, Created)
 DT 01-JUN-2001 (TrEMBLrel. 17, Last sequence update)
 DT 01-JUN-2001 (TrEMBLrel. 17, Last annotation update)
 DE 2'-5' OLIGOADENYLATE SYNTHETASE-LIKE.
 GN OASL.
 OS Mus musculus (Mouse).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
 OX NCBI_TaxID=10090;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN=C57BL/6J; TISSUE=TONGUE;
 RX MEDLINE=21085660; PubMed=11217851;
 RA Kawai J., Shinagawa A., Shibata K., Yoshino M., Itoh M., Ishii Y.,
 RA Arakawa T., Hara A., Fukunishi Y., Konno H., Adachi J., Fukuda S.,
 RA Aizawa K., Izawa M., Nishi K., Kiyosawa H., Kondo S., Yamanaka I.,
 RA Saito T., Okazaki Y., Gojobori T., Bono H., Kasukawa T., Saito R.,
 RA Kadota K., Matsuda H.A., Ashburner M., Batalov S., Casavant T.,

RA Fleischmann W., Gaasterland T., Gissi C., King B., Kochiwa H.,
 RA Kuehl P., Lewis S., Matsuo Y., Mikaido I., Pesole G., Quackenbush J.,
 RA Schriml L.M., Staubli F., Suzuki R., Tomita M., Wagner L., Washio T.,
 RA Sakai K., Okado T., Furuno M., Aono H., Baldarelli R., Barsh G.,
 RA Blake J., Boffelli D., Bojuna N., Carninci P., de Bonaldo M.F.,
 RA Brownstein M.J., Bult C., Fletcher C., Fujita M., Gariboldi M.,
 RA Gustincich S., Hill D., Hoimann M., Hume D.A., Kaniya M., Lee N.H.,
 RA Lyons P., Marchionni L., Mashima J., Mazzarelli J., Mombaerts P.,
 RA Nordone P., Ring B., Ringwald M., Rodriguez I., Sakamoto N.,
 RA Sasaki H., Sato K., Schoenbach C., Seya T., Shibata Y., Storch K.-F.,
 RA Suzuki H., Toyooka K., Wang K.H., Weitz C., Whittaker C., Wilming L.,
 RA Wyshaw-Boris A., Yoshida K., Hasegawa Y., Kawaji H., Kohtsuki S.,
 RA Hayashizaki Y.,
 RA "Functional annotation of a full-length mouse cDNA collection."
 RL Nature 403:685-690(2001).
 DR EMBL; AK010034; BAB26655.1; -.
 DR MGD; MGI:1344390; Oasl.
 DR InterPro; IPR001797; 25A_synth.
 DR InterPro; IPR001201; PAP_25A_core.
 DR InterPro; IPR000626; Ubiquitin.
 DR Pfam; PF00240; ubiquitin; 1.
 DR SMART; SM00213; Ubq; 1.
 DR PROSITE; PS00833; 25A_SYNTH_2; UNKNOWN_1.
 DR PROSITE; PS0152; 25A_SYNTH_3; 1.
 DR PROSITE; PS50053; UBIQUITIN_2; 1.
 SQ SEQUENCE 508 AA; 58767 MW; 9552B4540CC801A0 CRC64;

Query Match 85.4%; Score 35; DB 11; Length 508;
 Best Local Similarity 71.4%; Pred. No. 12;
 Matches 5; Conservative 2; Mismatches 0; Indels 0; Gaps 0;

QY 1 KKRIMHC 7
 |||::||
 DB 109 KKRILHC 115

RESULT 9
 Q9QXV3 PRELIMINARY; PRT; 279 AA.
 AC Q9QXV3; Q9QXV4; Q9QUP8; Q9QZK3;
 DT 01-MAY-2000 (TRENBLrel. 13, Created)
 DT 01-MAY-2000 (TRENBLrel. 13, Last sequence update)
 DT 01-JUN-2001 (TRENBLrel. 17, Last annotation update)
 DE INGI PROTEIN.
 GN INGI.
 OS Mus musculus (Mouse).
 CC Eukaryota; Chordata; Craniata; Vertebrata; Euteleostomi;
 CC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Mus.
 OX NCBI_TaxID=10090;
 [1]
 RP SEQUENCE FROM N.A., AND CHARACTERIZATION.
 RC STRAIN=129/SVJ; TISSUE=BRAIN, EMBRYONIC FIBROBLAST, AND SPLEEN;
 RX MEDLINE=20011419; PubMed=10542254;
 RA Zaremski M., Hill J.E., Kwek S.S.S., Grigorian I.A., Gurova K.V.,
 RA Garkavtsev I.V., Diatchenko L., Koonin E.V., Gudkov A.V.;
 RT "Structure and regulation of the mouse Ingi gene. Three alternative
 RT transcripts encode two PHD finger proteins that have opposite effects
 RT on p53 function."
 RL J. Biol. Chem. 274:32172-32181(1999).
 RN [2]
 RP SEQUENCE FROM N.A. (ISOFORM 1).
 RA Rancourt D., Garkavtsev I.;
 RT "Structural organization and expression pattern of the murine Ingi
 RT gene."
 RL Submitted (MAY-1999) to the EMBL/GenBank/DBJ databases.
 CC -!- FUNCTION: ISOFORM 1 INHIBITS P53-DEPENDENT TRANSCRIPTIONAL
 CC ACTIVATION AND MAY FUNCTION AS AN ONCOPROTEIN. ISOFORM 2 ACTS AS A
 CC NEGATIVE GROWTH REGULATOR BY COOPERATING WITH P53 IN
 CC TRANSCRIPTIONAL ACTIVATION OF P53-RESPONSIVE GENES AND MAY ACT AS
 CC A TUMOR SUPPRESSOR.
 CC -!- SUBCELLULAR LOCATION: NUCLEAR (BY SIMILARITY).
 CC -!- ALTERNATIVE PRODUCTS: 2 ISOFORMS; 1 (SHOWN HERE) AND 2; ARE

CC PRODUCED BY ALTERNATIVE SPLICING.
 CC -!- TISSUE SPECIFICITY: IN THE ADULT, WIDELY EXPRESSED WITH HIGHEST
 CC LEVELS IN THYMUS AND TESTIS. EXPRESSED THROUGHOUT THE WHOLE EMBRYO
 CC AT ALL STAGES OF DEVELOPMENT EXAMINED. AT DAY 10, HIGHEST
 CC EXPRESSION IS FOUND IN THE YOLK SAC WHILE AT DAY 16 AND 18, HIGHER
 CC LEVELS ARE FOUND IN INNER COMPARTMENTS OF BONE.
 CC -!- DEVELOPMENTAL STAGE: IN THE EMBRYO, HIGHEST EXPRESSION OF ISOFORM
 CC 1 IS FOUND AT DAY 11 WHILE HIGHEST EXPRESSION OF ISOFORM 2 IS
 CC FOUND AT DAY 7.
 CC -!- SIMILARITY: CONTAINS 1 PHD-FINGER DOMAIN.
 DR EMBL; ARL177753; AAF16911.1; -.
 DR EMBL; ARL177755; AAF16908.1; -.
 DR EMBL; ARL177756; AAF16909.1; -.
 DR EMBL; ARL177757; AAF16910.1; -.
 DR EMBL; ARL49820; AAF09183.1; -.
 DR MGD; MGI:1349481; Ingi.
 DR InterPro; IPR001965; PHD.
 DR Pfam; PF00628; PHD; 1.
 DR SMART; SM00249; PHD; 1.
 KW Oncogene; Anti-oncogene; Alternative splicing.
 FT DOMAIN 210 259 PHD-FINGER.
 FT VARSPLIC 1 94 MISSING (IN ISOFORM 2).
 FT CONFLICT 203 203 L -> F (IN REF. 2).
 SQ SEQUENCE 279 AA; 32109 MW; 6765C984EEF179F4 CRC64;

Query Match 82.9%; Score 34; DB 11; Length 279;
 Best Local Similarity 57.1%; Pred. No. 12;
 Matches 4; Conservative 3; Mismatches 0; Indels 0; Gaps 0;

QY 1 KKRIMHC 7
 |||::||
 DB 67 KRRVLHC 73

RESULT 10
 Q9M149 PRELIMINARY; PRT; 401 AA.
 AC Q9M149;
 DT 01-OCT-2000 (TRENBLrel. 15, Created)
 DT 01-OCT-2000 (TRENBLrel. 15, Last sequence update)
 DT 01-JUN-2001 (TRENBLrel. 17, Last annotation update)
 DE PUTATIVE PHOSPHATIDYLINOSITOL KINASE.
 GN AT4G01190.
 OS Arabidopsis thaliana (Mouse-ear cress).
 CC Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
 CC Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots; Rosidae;
 CC eurosids II; Brassicales; Brassicaceae; Arabidopsis.
 OX NCBI_TaxID=3702;
 [1]
 RP SEQUENCE FROM N.A.
 RA Lamar B., Stoneking T., Stumpf J., Mewes H.W., Lemcke K.,
 RA Mayer K.F.X.;
 RL Submitted (MAR-2000) to the EMBL/GenBank/DBJ databases.
 RN [2]
 RP SEQUENCE FROM N.A.
 RA EU Arabidopsis sequencing project;
 RL Submitted (MAR-2000) to the EMBL/GenBank/DBJ databases.
 DR EMBL; AL161491; CAB80928.1; -.
 DR InterPro; IPR002498; PIP5K.
 DR Pfam; PF01504; PIP5K; 1.
 DR SMART; SM00330; PIPKC; 1.
 KW Kinase.
 SQ SEQUENCE 401 AA; 45658 MW; 8A12D10DA2DEB4CA CRC64;

Query Match 82.9%; Score 34; DB 10; Length 401;
 Best Local Similarity 85.7%; Pred. No. 16;
 Matches 6; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 KKRIMHC 7
 |||||
 DB 353 KKRIEHC 359

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RESULT 11
004613 PRELIMINARY; PRT: 431 AA.
ID Q04613
AC Q04613;
DT 01-JUL-1997 (TREMBlrel. 04, Created)
DT 01-JUL-1997 (TREMBlrel. 04, Last sequence update)
DT 01-JUN-2001 (TREMBlrel. 17, Last sequence update)
DE SIMILAR TO PHOSPHATIDYLINOSITOL-4-PHOSPHATE -KINASE TYPE II.
GN A.IG002N01.9.
OS Arabidopsis thaliana (Mouse-ear cross).
OC Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
OC Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots; Rosidae;
OC eurosids II; Brassicales; Brassicaceae; Arabidopsids.
OX NCBI_TaxID=3702;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=CV. COLUMBIA;
RA Scheet P., Maggi L.;
RL Submitted (JUN-1997) to the EMBL/GenBank/DBJ databases.
RN [2]
RP SEQUENCE FROM N.A.
RC STRAIN=CV. COLUMBIA;
RA Waterston R.;
RL Submitted (JUN-1997) to the EMBL/GenBank/DBJ databases.
DR EMBL: AF007269; AAB61030.1; -.
DR Mendel; 17574; Arath;2860;17574.
DR InterPro: IPR002498; PIP5K.
DR Pfam: PF01504; PIP5K; 1.
DR SMART: SM00330; PIP5K; 1.
SQ SEQUENCE 431 AA; 49356 MW; 07A53F23BDDDD942B CRC64;

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Query Match 82.9%; Score 34; DB 10; Length 431;
Best Local Similarity 85.7%; Pred. No. 17;
Matches 6; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

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QY 1 KKRIMHC 7
Db 383 KKRIMHC 389

RESULT 12
Q03949 PRELIMINARY; PRT: 189 AA.
ID Q03949
AC Q03949;
DT 01-NOV-1996 (TREMBlrel. 01, Created)
DT 01-NOV-1996 (TREMBlrel. 01, Last sequence update)
DT 01-NOV-1998 (TREMBlrel. 08, Last annotation update)
DE INSERTION ELEMENT IS895 HYPOTHETICAL 21.9 KDA PROTEIN (ORFI).
OS Anabaena sp. (strain PCC 7120).
OC Bacteria; Cyanobacteria; Nostocales; Nostocaceae; Nostoc.
OX NCBI_TaxID=103690;
RN [1]
RP SEQUENCE FROM N.A.
RC MEDLINE=91358370; PubMed=1653219;
RX Alam J., Vrba J.M., Cai Y., Martin J.A., Weislo L.J., Curtis S.E.;
RT "Characterization of the IS895 family of insertion sequences from the
RT cyanobacterium Anabaena sp. strain PCC 7120.";
RL J. Bacteriol. 173:5778-5783 (1991).
DR EMBL: M67475; AAA98138.1; -.
DR FIr; A38117; A38117.
KW Hypothetical protein; Transposable element.
SQ SEQUENCE 189 AA; 21937 MW; B873A342856C2103 CRC64;

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Query Match 80.5%; Score 33; DB 2; Length 189;
Best Local Similarity 71.4%; Pred. No. 14;
Matches 5; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

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```

QY 1 KKRIMHC 7
|||||

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Db 171 KKRLKHC 177

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RESULT 13
Q9AKS0 PRELIMINARY; PRT: 267 AA.
ID Q9AKS0
AC Q9AKS0;
DT 01-JUN-2001 (TREMBlrel. 17, Created)
DT 01-JUN-2001 (TREMBlrel. 17, Last sequence update)
DT 01-JUN-2001 (TREMBlrel. 17, Last annotation update)
DE TATC PROTEIN.
GN TATC.
OS Pseudomonas stutzeri (Pseudomonas perfectomarina).
OC Bacteria; Proteobacteria; gamma subdivision; Pseudomonadaceae;
OC Pseudomonas.
OX NCBI_TaxID=316;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=ZOBELL ATCC14405;
RX MEDLINE=21101859; PubMed=11160097;
RA Heikkilae M.P., Honisch U., Wunsch P., Zumft W.G.;
RT "Role of the Tat transport system in nitrous oxide reductase
RT translocation and cytochrome cdl biosynthesis in Pseudomonas
RT stutzeri".
RL J. Bacteriol. 183:1663-1671 (2001).
RL EMBL: AJ299712; CAC29149.1; -.
SQ SEQUENCE 267 AA; 29439 MW; 4369EB5E38BCD736 CRC64;

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Query Match 80.5%; Score 33; DB 2; Length 267;
Best Local Similarity 57.1%; Pred. No. 19;
Matches 4; Conservative 3; Mismatches 0; Indels 0; Gaps 0;

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QY 1 KKRIMHC 7
|||||
Db 21 KKRLKHC 27

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RESULT 14
Q9ST63 PRELIMINARY; PRT: 495 AA.
ID Q9ST63
AC Q9ST63;
DT 01-MAY-2000 (TREMBlrel. 13, Created)
DT 01-MAY-2000 (TREMBlrel. 13, Last sequence update)
DT 01-JUN-2001 (TREMBlrel. 17, Last annotation update)
DE PUTATIVE INTERNAL ROTENONE-INSENSITIVE NADH DEHYDROGENASE.
GN NDA1.
OS Solanum tuberosum (Potato).
OC Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
OC Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots;
OC Asteridae; euasterids I; Solanales; Solanaceae; Solanum.
OX NCBI_TaxID=4113;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=CV. DESIREE; TISSUE=LEAF;
RA Rasmussen A.G., Svensson A., Knoop V., Grohmann L., Brennicke A.;
RT "Homologues of yeast and bacterial rotenone-insensitive NADH
RT dehydrogenases in higher eukaryotes: two enzymes are present in potato
RT mitochondria.";
RL Submitted (AUG-1999) to the EMBL/GenBank/DBJ databases.
CC -!- COFACTOR: FAD (BY SIMILARITY).
CC -!- SIMILARITY: TO PYRIDINE NUCLEOTIDE-DISULPHIDE OXIDOREDUCTASES
CC CLASS=I.
DR EMBL: AJ245861; CAB52796.1; -.
DR InterPro: IPR001327; FAD_pyr_redox.
DR Pfam: PF00070; pyr_redox; 1.
DR FAD: Flavoprotein; NAD: Oxidoreductase; Redox-active center.
KW SEQUENCE, 495 AA; 54902 MW; 6AFFC807BEB01340 CRC64;

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Query Match 80.5%; Score 33; DB 10; Length 495;
Best Local Similarity 57.1%; Pred. No. 32;
Matches 4; Conservative 3; Mismatches 0; Indels 0; Gaps 0;

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QY 1 KKRIMHC 7
|.:|.:|
Db 224 KRLLHC 230

RESULT 15

OB0874
ID OB0874 PRELIMINARY; PRT: 508 AA.
AC OB0874;
DT 01-NOV-1998 (TReMBLrel. 08, Created)
DT 01-NOV-1998 (TReMBLrel. 08, Last sequence update)
DE 01-JUN-2001 (TReMBLrel. 17, Last annotation update)
DE PUTATIVE UBIQUINONE REDUCTASE.
GN F23F1.9.
OS Arabidopsis thaliana (Mouse-ear cress).
OC Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
OC Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots; Rosidae;
OC eurosids II; Brassicales; Brassicaceae; Arabidopsis.
OX NCBI_TaxID=3702;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=CV. COLUMBIA;
RA Rounsley S.D., Lin X., Ketchum K.A., Crosby M.L., Brandon R.C.,
RA Sykes S.M., Mason T.M., Kerlavage A.R., Adams M.D.,
RA Somerville C.R., Venter J.C.;
RT "Arabidopsis thaliana chromosome II BAC F23F1 genomic sequence."
RL Submitted (AUG-1998) to the EMBL/GenBank/DBJ databases.
CC -!- COFACTOR: FAD (BY SIMILARITY).
CC -!- SIMILARITY: TO PYRIDINE NUCLEOTIDE-DISULPHIDE OXIDOREDUCTASES
CC CLASS-I.
DR EMBL; AC004680; AAC31853.1; .
DR InterPro: IPR001327; FAD_pyr_redox.
DR Pfam: PF00070; pyr_redox; 1.
KW FAD; Flavoprotein; Oxidoreductase; Redox-active center.
SQ SEQUENCE 508 AA; 56503 MW; 266A434E702A0C27 CRC64;

Query Match 80.5%; Score 33; DB 10; Length 508;
Best Local Similarity 57.1%; Pred. No. 33;
Matches 4; Conservative 3; Mismatches 0; Indels 0; Gaps 0;

QY 1 KKRIMHC 7
|.:|.:|
Db 236 KRLLHC 242

Search completed: February 12, 2002, 12:03:25
Job time: 816 sec

GenCore version 4.5
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OM protein - protein search, using sw model

Run on: February 12, 2002, 12:04:01 ; Search time 30.28 Seconds
(without alignments)
8.476 Million cell updates/sec

Title: US-09-606-129A-18

Perfect score: 41

Sequence: 1 KRRIMHC 7

Scoring table: BLOSUM62

Gapop 10.0 , Gapext 0.5

Searched: 100059 segs, 36664827 residues

Total number of hits satisfying chosen parameters: 100059

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database : SwissProt_39:**

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	41	100.0	295	1 BIEA_RAT	P46844 rattus norv
2	38	92.7	236	1 BIEA_HUMAN	P53004 homo sapien
3	35	85.4	473	1 OASL_MOUSE	Q92312 mus musculu
4	33	80.5	925	1 PIP1_YEAST	P40020 saccharomyc
5	32	78.0	140	1 FKBP2_MOUSE	P45878 mus musculu
6	32	78.0	141	1 FKBP2_HUMAN	P26885 homo sapien
7	32	78.0	328	1 HYBA_ECOLI	P37179 escherichia
8	32	78.0	528	1 DYR2_HUMAN	Q92630 homo sapien
9	32	78.0	571	1 AMPL_LICES	Q10712 lycopersico
10	32	78.0	573	1 AMPL_SOLTU	P31427 solanum tub
11	31	75.6	105	1 RL44_TRYBB	P17843 trypanosoma
12	31	75.6	279	1 Y091_NPVOP	O10341 orgyia pseu
13	31	75.6	323	1 CYCH_XENLA	P51947 xenopus lae
14	31	75.6	451	1 SUN_HAEIN	P44788 haemophilus
15	31	75.6	768	1 CUL3_HUMAN	Q13618 homo sapien
16	31	75.6	768	1 CUL3_MOUSE	Q9JJV5 mus musculu
17	31	75.6	861	1 UL52_HSV7J	P52468 human herpe
18	30	73.2	108	1 Y209_METJA	Q60271 methanococc
19	30	73.2	4128	1 PRKD_MOUSE	P97313 mus musculu
20	29	70.7	38	1 RL36_THEMA	Q9X116 thermotoga
21	29	70.7	142	1 SECB_BUCAI	P57161 buchnera ap
22	29	70.7	247	1 YHP5_YEAST	P38908 saccharomyc
23	29	70.7	257	1 YZGL_CAEEL	P55266 caenorhabdi
24	29	70.7	300	1 RANT_BPF22	P03037 bacterioph
25	29	70.7	359	1 ODPB_RAT	P49432 rattus norv
26	29	70.7	423	1 SHP1_YEAST	P34223 saccharomyc
27	29	70.7	560	1 YD2H_SCHPO	Q10264 schizosacch
28	29	70.7	588	1 DYR3_HUMAN	Q43781 homo sapien
29	29	70.7	632	1 AFUB_HAEIN	Q57341 haemophilus
30	29	70.7	670	1 REP_HAEIN	P44604 haemophilus
31	29	70.7	918	1 YK62_CAEEL	P34341 caenorhabdi
32	29	70.7	1059	1 CAPU_DROME	Q24120 drosophila
33	29	70.7	1790	1 SEPA_EWENI	P78621 emericeila

ALIGNMENTS

RESULT 1

ID	BIEA_RAT	STANDARD;	PRT;	295 AA.
AC	P46844;			
DT	01-NOV-1995 (Rel. 32, Created)			
DT	01-NOV-1995 (Rel. 32, Last sequence update)			
DT	15-JUL-1999 (Rel. 38, Last annotation update)			
DE	BILIVERDIN REDUCTASE A PRECURSOR (EC 1.3.1.24) (BILIVERDIN-IX ALPHA-REDUCTASE).			
DE	REDUCTASE).			
GN	BLVRA OR BLVR.			
OS	Rattus norvegicus (Rat).			
OC	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;			
OC	Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Rattus.			
OX	NCBI_TaxID=10116;			
RN	[1]			
RP	SEQUENCE FROM N.A., AND PARTIAL SEQUENCE.			
RC	TISSUE=Kidney; PubMed=1371282;			
RX	MEDLINE=92156147; PubMed=1371282;			
RA	Fakhrai H., Maines M.D.;			
RT	"Expression and characterization of a cDNA for rat kidney biliverdin reductase. Evidence suggesting the liver and kidney enzymes are the same transcript product."			
RT	same transcript product."			
RL	J. Biol. Chem. 267:4023-4029(1992).			
RN	[2]			
RP	MUTAGENESIS.			
RC	MEDLINE=94291657; PubMed=8020496;			
RA	McCoubrey W.K. Jr., Maines M.D.;			
RT	"Site-directed mutagenesis of cysteine residues in biliverdin reductase. Roles in substrate and cofactor binding."			
RL	Eur. J. Biochem. 222:597-603(1994).			
CC	!- FUNCTION: CONVERTS BILIVERDIN TO BILIRUBIN. DISPLAYS TWO DISTINCT PH OPTIMA USING A DIFFERENT COFACTOR AT EACH PH: NADH AT THE LOWER PH 6.7-6.9 RANGE AND NADPH AT PH 8.5-8.7. NADPH, HOWEVER, IS THE PROBABLE COFACTOR IN BIOLOGICAL SYSTEMS.			
CC	!- CATALYTIC ACTIVITY: BILIRUBIN + NAD(P)(+) = BILIVERDIN + NAD(P)H.			
CC	!- COFACTOR: BINDS ONE ZINC ION.			
CC	!- PATHWAY: FINAL STEP IN HEME METABOLISM.			
CC	!- SUBUNIT: MONOMER (BY SIMILARITY).			
CC	!- SUBCELLULAR LOCATION: CYTOPLASMIC.			
CC	!- SIMILARITY: TO E.COLI YHHX.			
CC	-----			
CC	This SWISS-PROT entry is copyright. It is produced through a collaboration between the Swiss Institute of Bioinformatics and the EMBL outstation - the European Bioinformatics Institute. There are no restrictions on its use by non-profit institutions as long as its content is in no way modified and this statement is not removed. Usage by and for commercial entities requires a license agreement (See http://www.isb-sib.ch/announce/ or send an email to license@isb-sib.ch).			
CC	-----			
DR	EMBL; M81681; AAA40830.1; -			
DR	InterPro: IPR000583; GFO_IDH_MCCA.			
DR	Pfam; PF01408; GFO_IDH_MCCA; 1.			
KW	Oxidoreductase; NAD; NADP; Zinc.			
FT	PROPEP 1 2			
FT	CHAIN 3 295 BILIVERDIN REDUCTASE A.			
FT	DOMAIN 11 16 POLY-VAL.			

P39740 bacillus su
Q92CW5 rickettsia
P49766 mus musculu
P05449 rhodospheudo
P5910 bacillus su
Q53479 methanobact
P24442 dictyosteli
P16894 dictyosteli
O52199 mycobacteri
P51956 homo sapien
P98119 desmodus ro
P15638 desmodus ro

34 28 68.3 113 1 FLIT_BACSU
35 28 68.3 172 1 IPVR_RICPR
36 28 68.3 188 1 VRGB_MOUSE
37 28 68.3 249 1 YAT6_RHOBL
38 28 68.3 313 1 YQAM_BACSU
39 28 68.3 324 1 IDSA_METTM
40 28 68.3 335 1 UL16_HSV69
41 28 68.3 356 1 GRAL_DICDI
42 28 68.3 428 1 DGTP_MYCSM
43 28 68.3 459 1 NEK3_HUMAN
44 28 68.3 477 1 URT1_DESRO
45 28 68.3 477 1 URT2_DESRO

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FT METAL 279 279 ZINC (POTENTIAL).
FT METAL 280 280 ZINC (POTENTIAL).
FT METAL 291 291 ZINC (POTENTIAL).
FT METAL 292 292 ZINC (POTENTIAL).
FT MUTAGEN 73 73 C->A: LOSS OF ACTIVITY.
FT MUTAGEN 280 280 C->A: REDUCED ACTIVITY.
FT MUTAGEN 291 291 C->A: REDUCED ACTIVITY.
SQ SEQUENCE 295 AA: 33565 MW: 219C8EA96C150589 CRC64;

Query Match 100.0%; Score 41; DB 1; Length 295;
Best Local Similarity 100.0%; Pred. No. 0.13;
Matches 7; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 KKRIMHC 7
DB 274 KKRIMHC 280

RESULT 2
BIEA_HUMAN STANDARD; PRT; 296 AA.
AC P53004;
DT 01-OCT-1996 (Rel. 34, Created)
DT 01-OCT-1996 (Rel. 34, Last sequence update)
DT 20-AUG-2001 (Rel. 40, Last annotation update)
DE BILIVERDIN REDUCTASE A PRECURSOR (EC 1.3.1.24) (BILIVERDIN-IX ALPHA-
REDUCTASE).
GN BLVRA OR BLVR OR BVR.
OS Homo sapiens (Human).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
OX NCBI_TaxID=9606;
RN [1]
RP SEQUENCE FROM N.A.
RC TISSUE=Placenta;
RX MEDLINE=96202961; PubMed=8631357;
RA Maines M.D., Polevoda B.V., Huang T.-J., McCoubrey W.K. Jr.;
RT "Human biliverdin IXalpha reductase is a zinc-metalloprotein.
Characterization of purified and Escherichia coli expressed
enzymes.";
RL Eur. J. Biochem. 235:372-381(1996).
RN [2]
RP SEQUENCE FROM N.A.
RA Komuro A., Tobe T., Nakano Y., Yamaguchi T., Tomita M.;
RL Submitted (JAN-1996) to the EMBL/GenBank/DBJ databases.
RN [3]
RP SEQUENCE OF 1-117 FROM N.A.
RA Cordes M., Wollam C., Carter T.;
RL Submitted (JUN-1998) to the EMBL/GenBank/DBJ databases.
RN [4]
RP SEQUENCE OF 3-36; 48-74 AND 228-248.
RC TISSUE=Liver;
RX MEDLINE=93143333; PubMed=8424566;
RA Maines M.D., Trakshel G.M.;
RT "Purification and characterization of human biliverdin reductase.";
RL Arch. Biochem. Biophys. 300:320-326(1993).
RN [5]
RP SEQUENCE OF 3-22.
RC TISSUE=Liver;
RX MEDLINE=95014177; PubMed=7929092;
RA Yamaguchi T., Komoda Y., Nakajima H.;
RT "Biliverdin-IX alpha reductase and biliverdin-IX beta reductase from
human liver. Purification and characterization.";
RL J. Biol. Chem. 269:24343-24348(1994).
CC -!- FUNCTION: CONVERTS BILIVERDIN TO BILIRUBIN.
CC -!- CATALYTIC ACTIVITY: BILIRUBIN + NAD(P)(+) = BILIVERDIN + NAD(P)H.
CC -!- COFACTOR: BINDS ONE ZINC ION. HAS DUAL PH/COFACTOR (NADH, NADPH)
SPECIFICITY. USES NADH AT THE ACIDIC PH RANGE (6-6.7) AND NADPH AT
THE ALKALINE RANGE (8.5-8.7).
CC -!- PATHWAY: FINAL STEP IN HEME METABOLISM.
CC -!- SUBUNIT: MONOMER.
CC -!- SUBCELLULAR LOCATION: CYTOPLASMIC.

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CC -!- TISSUE SPECIFICITY: LIVER.
CC -!- SIMILARITY: TO E.COLI YHHX.
CC -----
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CC -----
DR EMBL; X93086; CAA63635.1;
DR EMBL; U34877; AAC35588.1;
DR EMBL; AC005189; AAC25526.1;
DR MIM; 109750;
DR InterPro; IPR000683; GFO_IDH_MOCA.
DR Pfam; PF01408; GFO_IDH_MOCA; 1.
KW Oxidoreductase; NAD; NADP; Zinc.
FT PROPEP 1 2
FT CHAIN 3 296 BILIVERDIN REDUCTASE A.
FT DOMAIN 11 16 POLY-VAL.
FT METAL 280 280 ZINC (POTENTIAL).
FT METAL 281 281 ZINC (POTENTIAL).
FT METAL 292 292 ZINC (POTENTIAL).
FT METAL 293 293 ZINC (POTENTIAL).
FT CONFLICT 3 3 A -> T (IN REF. 2).
FT CONFLICT 154 155 SD -> AG (IN REF. 2).
FT CONFLICT 160 160 D -> E (IN REF. 2).
SQ SEQUENCE 296 AA; 33488 MW; 0DFD3B86F4DFC0A CRC64;

Query Match 92.7%; Score 38; DB 1; Length 296;
Best Local Similarity 85.7%; Pred. No. 0.55;
Matches 6; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

QY 1 KKRIMHC 7
DB 275 KKRILHC 281

RESULT 3
OASL_MOUSE STANDARD; PRT; 473 AA.
AC Q9Z2F2;
DT 20-AUG-2001 (Rel. 40, Created)
DT 20-AUG-2001 (Rel. 40, Last sequence update)
DT 20-AUG-2001 (Rel. 40, Last annotation update)
DE 5A KDA 2'-5'-OLIGOADENYLATE SYNTHETASE LIKE PROTEIN (EC 2.7.7.-) (P54
OASL) (P54OASL) (M1204).
GN OASL.
OS Mus musculus (Mouse).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
OX NCBI_TaxID=10090;
RN [1]
RP SEQUENCE FROM N.A.
RC TISSUE=Spleen;
RX MEDLINE=99323969; PubMed=10395668;
RA Tiefenthaler M., Marksteiner R., Neyer S., Koch F., Hofer S.,
RA Schuler G., Nussenzweig M., Schneider R., Heufler C.;
RT "M1204, a novel 2',5'-oligoadenylate synthetase with a ubiquitin-like
extension, is induced during maturation of murine dendritic cells.";
RL J. Immunol. 163:760-765(1999).
CC -!- FUNCTION: MAY PLAY A ROLE IN MEDIATING RESISTANCE TO VIRUS
INFECTION. CONTROL OF CELL GROWTH, DIFFERENTIATION, AND APOPTOSIS.
CC -!- CATALYTIC ACTIVITY: BINDS DOUBLE-STRANDED RNA AND POLYMERIZES ATP
INTO PPP(A2'P5'A)N OLIGOMERS, WHICH ACTIVATE THE LATENT RNASE L
THAT, WHEN ACTIVATED, CLEAVES SINGLE-STRANDED RNAs.
CC -!- TISSUE SPECIFICITY: STRONGLY EXPRESSED IN SPLEEN DENDRITIC CELLS.
WHEREAS IN BONE MARROW-DERIVED DENDRITIC CELLS, THE AMOUNT
INCREASES DURING THE MATURATION PROCESS. EXPRESSED IN MANY ORGANS,
THE HIGHEST LEVELS BEING IN THYMUS, LUNG, AND BONE MARROW.
CC -!- SIMILARITY: BELONGS TO THE 2-5A SYNTHETASE FAMILY.

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CC -!- SIMILARITY: CONTAINS 1 UBIQUITIN-LIKE DOMAIN.
CC -!- CAUTION: THIS MAY NOT BE THE TRUE ORTHOLOG OF HUMAN OASL.
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CC or send an email to license@isb-sib.ch).
CC -----
CC EMBL: AF068835; AAD02818.1; -
CC DR MGD; MGI:1344390; Casl.
CC DR InterPro: IPR001797; 25A_synth.
CC DR InterPro: IPR001201; PAP_25A_core.
CC DR Pfam: PF00240; ubiquitin; 1.
CC DR PROSITE: PS00832; 25A_SYNTH_1; FALSE_NEG.
CC DR PROSITE: PS00833; 25A_SYNTH_2; 1.
CC DR PROSITE: PS0152; 25A_SYNTH_3; 1.
CC DR PROSITE: PS50053; UBIQUITIN_2; FALSE_NEG.
CC DR PROSITE: PS50053; UBIQUITIN_3; 1.
CC DR RNA-binding; Transferase; Nucleotidyltransferase.
CC KW DOMAIN 435 473 UBIQUITIN-LIKE.
CC FT SEQUENCE 473 AA; 54625 MW; 570E0E08A51C8460 CRC64;
CC -----
CC Query Match 85.4%; Score 35; DB 1; Length 473;
CC Best Local Similarity 71.4%; Pred. No. 3.8;
CC Matches 5; Conservative 2; Mismatches 0; Indels 0; Gaps 0;
CC -----
CC QY 1 KKRIMHC 7
CC Db 109 KKRLIHC 115
CC -----
CC RESULT 4
CC ID P1P1_YEAST STANDARD; PRT; 925 AA.
CC AC P40020; 1995 (Rel. 31, Created)
CC DT 01-FEB-1997 (Rel. 35, Last sequence update)
CC DT 01-NOV-1997 (Rel. 35, Last annotation update)
CC DE POLYMERASE-INTERACTING PROTEIN 1 (FACTOR INTERACTING WITH REF).
CC GN P1P1 OR FIRL OR YER032W.
CC OS Saccharomyces cerevisiae (Baker's yeast).
CC OC Eukaryota; Fungi; Ascomycota; Saccharomycotina; Saccharomycetes;
CC OC Saccharomycetales; Saccharomycetaceae; Saccharomycetes.
CC OX NCBI_TaxID=4932;
CC RN [1]
CC RP SEQUENCE FROM N.A.
CC RC STRAIN=W303;
CC RA del Olmo M., Gross S., Moore C.L.;
CC RL Submitted (FEB-1997) to the EMBL/GenBank/DBJ databases.
CC RN [2]
CC RP SEQUENCE FROM N.A.
CC RC STRAIN=S288C / AB972;
CC RA Dietrich F.S., Mulligan J.T., Hennessey K.M., Allen E., Araujo R.,
CC RA Aviles E., Berno A., Brennan T., Carpenter J., Chen E., Cherry J.M.,
CC RA Chung E., Duncan M., Guzman E., Hartzell G., Hunnicke-Smith S.,
CC RA Hyman R., Kayser A., Komp C., Lashkari D., Lew H., Lin D.,
CC RA Moresdale D., Nakahara K., Namath A., Norgren R., Oeiner P., Oh C.,
CC RA Patel F.X., Roberts D., Sehl P., Schramm S., Shogren T., Smith V.,
CC RA Taylor P., Wei Y., Yelton M., Botstein D., Davis R.W.;
CC RL Submitted (DEC-1994) to the EMBL/GenBank/DBJ databases.
CC RN [3]
CC RP CHARACTERIZATION.
CC RN MEDLINE=97339480; PubMed=9196079;
CC RA Russnak R., Pereira S., Platt T.;
CC FT "RNA binding analysis of yeast REF2 and its two-hybrid interaction
CC with a new gene product, FIRL."
CC RL Gene Expr. 6:241-258(1996).
CC CC -!- FUNCTION: INTERACTS WITH POLY(A) POLYMERASE AND WITH REF2.
CC -----

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CC -----
CC EMBL: U17262; AAB46625.1; -
CC DR EMBL; U18778; AAB64565.1; -
CC DR SGD; S0000834; FIR1 663
CC FT CONFLICT 663
CC SQ SEQUENCE 925 AA; 104701 MW; 707D9839EE31322B CRC64;
CC -----
CC Query Match 80.5%; Score 33; DB 1; Length 925;
CC Best Local Similarity 71.4%; Pred. No. 19;
CC Matches 5; Conservative 1; Mismatches 1; Indels 0; Gaps 0;
CC -----
CC QY 1 KKRIMHC 7
CC Db 841 KKRLSHC 847
CC -----
CC RESULT 5
CC ID FKX2_MOUSE STANDARD; PRT; 140 AA.
CC AC P45878;
CC DT 01-NOV-1995 (Rel. 32, Created)
CC DT 01-NOV-1995 (Rel. 32, Last sequence update)
CC DT 15-JUL-1999 (Rel. 38, Last annotation update)
CC DE FK506-BINDING PROTEIN PRECURSOR (FKBP-13) (PEPTIDYL-PROLYL CIS-TRANS
CC ISOMERASE) (PPIASE) (EC 5.2.1.8).
CC GN FKBP2 OR FKBP13.
CC OS Mus musculus (Mouse).
CC OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
CC OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
CC OX NCBI_TaxID=10090;
CC RN [1]
CC RP SEQUENCE FROM N.A.
CC RC STRAIN=129/SVJ; TISSUE=Liver;
CC RX MEDLINE=94085790; PubMed=7505249;
CC RA Hendrickson B.A., Zhang W., Craig R.J., Jin Y.J., Blier R.E.,
CC RA Burakoff S.J., Dilella A.G.;
CC RT "Structural organization of the genes encoding human and murine
CC FK506-binding protein (FKBP) 13 and comparison to FKBP1."
CC RL Gene 134:271-275(1993).
CC CC -!- FUNCTION: PPIASES ACCELERATE THE FOLDING OF PROTEINS.
CC CC -!- CATALYTIC ACTIVITY: CIS-TRANS ISOMERIZATION OF PROLINE IMIDIC
CC CC -!- ENZYME REGULATION: INHIBITED BY BOTH FK506 AND RAPAMYCIN.
CC CC -!- SUBCELLULAR LOCATION: ENDOPLASMIC RETICULUM LUMEN, MEMBRANE
CC CC ASSOCIATED (PROBABLE).
CC CC -!- SIMILARITY: BELONGS TO THE FKBP-TYPE PPIASE FAMILY.
CC -----
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CC or send an email to license@isb-sib.ch).
CC -----
CC EMBL: M77831; AAA37631.1; -
CC DR HSP; P20081; 1YAT.
CC DR MGD; MGI:95542; FKBP2.
CC DR InterPro: IPR001179; FKBP_PPIase.
CC DR Pfam: PF00254; FKBP; 1.
CC DR PROSITE: PS00453; FKBP_PPIASE_1; 1.
CC DR PROSITE: PS00454; FKBP_PPIASE_2; 1.
CC DR PROSITE: PS00509; FKBP_PPIASE_3; 1.
CC KW Isomerase; Rotamase; Signal; Endoplasmic reticulum.
CC FT SIGNAL 1 22
CC -----

```

FT CHAIN 23 140 FK506-BINDING PROTEIN.
FT SITE 137 140 PREVENT SECRETION FROM ER (POTENTIAL).
SQ SEQUENCE 140 AA: 15344 MW: F4E7FCC7766A0416 CRC64;

Query Match 78.0%; Score 32; DB 1; Length 140;
Best Local Similarity 71.4%; Pred. NO. 4.8;
Matches 5; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

QY 1 KKRIMHC 7
Db 34 KKRVDHC 40

RESULT 6
FXB2_HUMAN
ID FXB2_HUMAN STANDARD: PRT; 141 AA.
AC P26885;
DT 01-AUG-1992 (Rel. 23, Created)
DT 01-AUG-1992 (Rel. 23, Last sequence update)
DT 15-JUL-1999 (Rel. 39, Last annotation update)
DE FK506-BINDING PROTEIN PRECURSOR (FKBP-13) (PEPTIDYL-PROLYL CIS-TRANS
DE ISOMERASE) (PPIASE) (EC 5.2.1.8).
GN FKBP2 OR FKBP13.
OS Homo sapiens (Human).
OC Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
OX NCBI_TaxID=9606;
RN [1]
RP TISSUE-Colon carcinoma;
RX MEDLINE=91319747; PubMed=1713687;
RA Jin Y.-J., Alberts M.W., Lane W.S., Bierer B.E., Schreiber S.L.,
RA Burakoff S.J.;
RT "Molecular cloning of a membrane-associated human FK506- and
RT rapamycin-binding protein, FKBP-13."
RL Proc. Natl. Acad. Sci. U.S.A. 88:6677-6681(1991).
RN [2]
SEQUENCE FROM N.A.
RX MEDLINE=93112052; PubMed=1281998;
RA Dilella A.G., Hawkins A., Craig R.J., Schreiber S.L., Griffin C.A.;
RT "Chromosomal band assignments of the genes encoding human FKBP12 and
RT FKBP13."
RL Biochem. Biophys. Res. Commun. 189:819-823(1992).
CC -!- FUNCTION: PPIASES ACCELERATE THE FOLDING OF PROTEINS.
CC -!- CATALYTIC ACTIVITY: CIS-TRANS ISOMERIZATION OF PROLINE IMIDIC
CC PEPTIDE BONDS IN OLIGOPEPTIDES.
CC -!- ENZYME REGULATION: INHIBITED BY BOTH FK506 AND RAPAMYCIN.
CC -!- SUBCELLULAR LOCATION: ENDOPLASMIC RETICULUM LUMEN. MEMBRANE
CC ASSOCIATED (PROBABLE).
CC -!- TISSUE SPECIFICITY: T-CELLS AND THYMUS.
CC -!- SIMILARITY: BELONGS TO THE FKBP-TYPE PPIASE FAMILY.
-----
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-----
DR EMBL; M65128; AAA58473.1; .
DR EMBL; M75099; AAA36563.1; .
DR PIR; JC1365; JC1365.
DR HSP; P20071; 1FRT.
DR MIM; 186946; .
DR InterPro; IPR001179; FKBP_PPIase.
DR Pfam; PF00254; FKBP; 1.
DR PROSITE; PS00453; FKBP_PPIASE_1; 1.
DR PROSITE; PS00454; FKBP_PPIASE_2; 1.
DR PROSITE; PS00059; FKBP_PPIASE_3; 1.
KW Isomerase; Rotamase; Signal; Endoplasmic reticulum; Polymorphism.
FT SIGNAL 1 21

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FT CHAIN 22 141 FK506-BINDING PROTEIN.
FT SITE 138 141 PREVENT SECRETION FROM ER (POTENTIAL).
FT VARIANT 21 21 S -> TA.
FT VARIANT 24 24 /FTIG=VAR_006410.
FT VARIANT 24 24 T -> A.
FT VARIANT 96 96 /FTIG=VAR_006411.
FT VARIANT 96 96 Y -> C.
FT /FTIG=VAR_006412.
SQ SEQUENCE 141 AA: 15654 MW: 9F4751CA7D82D064 CRC64;

Query Match 78.0%; Score 32; DB 1; Length 141;
Best Local Similarity 71.4%; Pred. NO. 4.9;
Matches 5; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

QY 1 KKRIMHC 7
Db 35 KKRVDHC 41

RESULT 7
HYBA_ECOLI
ID HYBA_ECOLI STANDARD: PRT; 328 AA.
AC P37179;
DT 01-OCT-1994 (Rel. 30, Created)
DT 01-OCT-1994 (Rel. 30, Last sequence update)
DT 20-AUG-2001 (Rel. 40, Last annotation update)
DE HYDROGENASE-2 OPERON PROTEIN HYBA PRECURSOR.
GN HYBA OR B2996 OR Z4350 OR ECS3881.
OS Escherichia coli, and
OS Escherichia coli O157:H7.
OC Bacteria; Proteobacteria; gamma subdivision; Enterobacteriaceae;
OC Escherichia.
OX NCBI_TaxID=562, 83334;
RN [1]
SEQUENCE FROM N.A.
RX STRAIN=K12 / TGI;
RX MEDLINE=94292472; PubMed=8021236;
RA Menon N.K., Chatelus C.Y., Dervartanian M., Wendt J.C.,
RA Shannugam K.T., Peck H.D. Jr., Przybyla A.E.;
RT "Cloning, sequencing, and mutational analysis of the hyb operon
RT encoding Escherichia coli hydrogenase 2."
RL J. Bacteriol. 176:4416-4423(1994).
RN [2]
SEQUENCE FROM N.A.
RX STRAIN=K12 / MG1655;
RX MEDLINE=97426617; PubMed=9278503;
RA Blattner F.R., Plunkett G. III, Burland V., Perna N.T., Burland V.,
RA Riley M., Collado-Vides J., Glasner J.D., Rode C.K., Mayhew G.F.,
RA Gregor J., Davis N.W., Kirkpatrick H.A., Goeden M.A., Rose D.J.,
RA Mau B., Shao Y.;
RT "The complete genome sequence of Escherichia coli K-12."
RL Science 277:1453-1474(1997).
RN [3]
SEQUENCE FROM N.A.
RX STRAIN=O157:H7 / EDL933 / ATCC 700927;
RX MEDLINE=21074935; PubMed=11206551;
RA Perna N.T., Plunkett G. III, Burland V., Mau B., Glasner J.D.,
RA Rose D.J., Mayhew G.F., Evans P.S., Gregor J., Kirkpatrick H.A.,
RA Posfai G., Hackett J., Klink S., Boutin A., Shao Y., Miller L.,
RA Grobeck E.J., Davis N.W., Lim A., Dimalanta E.T., Potamousis K.,
RA Apodaca J., Anantharaman T.S., Lin J., Yen G., Schwartz D.C.,
RA Welch R.A., Blattner F.R.;
RT "Genome sequence of enterohaemorrhagic Escherichia coli O157:H7."
RL Nature 409:529-533(2001).
RN [4]
SEQUENCE FROM N.A.
RX STRAIN=O157:H7 / RIMD 0509952;
RX MEDLINE=21156231; PubMed=11258796;
RA Hayashi T., Makino K., Ohnishi M., Kurokawa K., Ishii K., Yokoyama K.,
RA Han C.-G., Ohtsubo E., Nakayama K., Murata T., Tanaka M., Tobe T.,
RA Iida T., Takami H., Honda T., Sasakawa C., Ogasawara N., Yasunaga T.,
RA Kuhara S., Shiba T., Hattori M., Shinagawa H.;

```

"Complete genome sequence of enterohemorrhagic Escherichia coli O157:H7 and genomic comparison with a laboratory strain K-12";
 RL DNA Res. 8:11-22(2001)
 CC -!- FUNCTION: PARTICIPATES IN THE PERIPLASMIC ELECTRON-TRANSFERRING
 CC ACTIVITY OF HYDROGENASE 2 DURING ITS CATALYTIC TURNOVER.
 CC -!- COFACTOR: BINDS 3 4FE-4S CLUSTERS AND A 3FE-4S CLUSTER.
 CC -!- SUBCELLULAR LOCATION: PERIPLASMIC.
 CC -!- SIMILARITY: THE IRON-SULFUR CENTERS ARE SIMILAR TO THOSE OF
 CC 'BACTERIAL-TYPE' 4FE-4S FERREDOXINS.
 CC -!- CAUTION: WAS ORIGINALLY (REF.1) THOUGHT TO BE THE SMALL SUBUNIT
 CC OF HYDROGENASE 2.
 CC
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 CC
 CC EMBL; U09177; AAA21589.1; -;
 CC EMBL; U28377; AAA69163.1; -;
 CC EMBL; AE000382; AAC76032.1; -;
 CC EMBL; AE005525; AAG58133.1; -;
 CC EMBL; AP002563; BAB37304.1; -;
 CC HSP; P55907; IXER.
 CC Ecogene; EGI1759; hybA.
 CC InterPro; IPR001450; 4FE4S_ferrdxin.
 CC Pfam; PF00037; fer4; 1.
 CC PROSITE; PS00198; 4FE4S_FERREDOXIN; 1.
 CC Oxidoreductase; Signal; Periplasmic; Iron-sulfur; 4Fe-4S; 3Fe-4S;
 KW Complete proteome.
 FT SIGNAL 1 27 POTENTIAL.
 FT CHAIN 28 328 HYDROGENASE-2 OPERON PROTEIN HYBA.
 FT METAL 47 47 IRON-SULFUR 1 (4FE-4S) (POTENTIAL).
 FT METAL 50 50 IRON-SULFUR 1 (4FE-4S) (POTENTIAL).
 FT METAL 53 53 IRON-SULFUR 1 (4FE-4S) (POTENTIAL).
 FT METAL 57 57 IRON-SULFUR 1 (4FE-4S) (POTENTIAL).
 FT METAL 112 112 IRON-SULFUR 2 (3FE-4S) (POTENTIAL).
 FT METAL 115 115 IRON-SULFUR 2 (3FE-4S) (POTENTIAL).
 FT METAL 120 120 IRON-SULFUR 2 (3FE-4S) (POTENTIAL).
 FT METAL 124 124 IRON-SULFUR 2 (3FE-4S) (POTENTIAL).
 FT METAL 145 145 IRON-SULFUR 3 (4FE-4S) (POTENTIAL).
 FT METAL 148 148 IRON-SULFUR 3 (4FE-4S) (POTENTIAL).
 FT METAL 151 151 IRON-SULFUR 3 (4FE-4S) (POTENTIAL).
 FT METAL 155 155 IRON-SULFUR 3 (4FE-4S) (POTENTIAL).
 FT METAL 174 174 IRON-SULFUR 4 (4FE-4S) (POTENTIAL).
 FT METAL 177 177 IRON-SULFUR 4 (4FE-4S) (POTENTIAL).
 FT METAL 193 193 IRON-SULFUR 4 (4FE-4S) (POTENTIAL).
 FT METAL 197 197 IRON-SULFUR 4 (4FE-4S) (POTENTIAL).
 SQ SEQUENCE 328 AA; 36003 MW; 77203A0F50F61662 CRC64;

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 CC Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
 CC NCBI_TaxID=9606;
 CC [1]
 CC SEQUENCE FROM N.A.
 CC RN
 CC TISSUE=Brain;
 CC MEDLINE=98421512; PubMed=9748265;
 CC Becker W., Weber Y., Wetzel K., Eimbter K., Tejedor F.J.,
 CC Joost H.-G.;
 CC "Sequence characteristics, subcellular localization, and substrate
 CC specificity of DYRK-related kinases, a novel family of dual
 CC specificity protein kinases";
 CC J. Biol. Chem. 273:25893-25902(1998).
 CC [2]
 CC SEQUENCE OF 320-528 FROM N.A.
 CC RN
 CC TISSUE=Placenta;
 CC Becker W., Joost H.-G.;
 CC Submitted (NOV-1996) to the EMBL/GenBank/DBJ databases
 CC EL
 CC -!- FUNCTION: IN VITRO; CAN PHOSPHORYLATE HISTONES H3 AND H2B ON SER
 CC AND THR RESIDUES. MAY BE INVOLVED IN THE REGULATION OF CELLULAR
 CC GROWTH AND/OR DEVELOPMENT.
 CC -!- SUBCELLULAR LOCATION: CYTOPLASMIC.
 CC -!- PTM: AUTOPHOSPHORYLATED ON TYR RESIDUES.
 CC -!- SIMILARITY: BELONGS TO THE SER/THR FAMILY OF PROTEIN KINASES.
 CC MNB/DYRK SUBFAMILY.
 CC
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 CC
 CC EMBL; Y13493; CAA73885.1; -;
 CC EMBL; Y09216; CAA70418.1; -;
 CC HSP; Q16539; 1WFC.
 CC MIN; 603496; -;
 CC InterPro; IPR000719; Euk_pkinase.
 CC InterPro; IPR002290; Ser_thr_kin_actsite.
 CC Pfam; PF00059; pkinase; 2.
 CC SMART; SM00220; S-TKC; 1.
 CC PROSITE; PS00107; PROTEIN_KINASE_ATP; 1.
 CC PROSITE; PS50011; PROTEIN_KINASE_DOM; 1.
 CC PROSITE; PS00108; PROTEIN_KINASE_ST; 1.
 CC Transferrase; Serine/threonine-protein kinase; Tyrosine-protein kinase;
 CC ATP-binding; Phosphorylation.
 CC DOMAIN 149 462 PROTEIN KINASE.
 CC NP_BIND 155 163 ATP (BY SIMILARITY).
 CC BINDING 178 178 ATP (BY SIMILARITY).
 CC ACT_SITE 275 275 BY SIMILARITY.
 SQ SEQUENCE 528 AA; 59714 MW; AF2C6822ED9522D7 CRC64;

Query Match 78.0%; Score 32; DB 1; Length 328;
 Best Local Similarity 71.4%; Pred. No. 11;
 Matches 5; Conservative 1; Mismatches 1; Indels 0; Gaps 0;
 QY 1 KKRIMHC 7
 DB 109 KKOCMHC 115
 RESULT 8
 ID_DYR2_HUMAN STANDARD; PRT; 528 AA.
 AC Q92630;
 DT 20-AUG-2001 (Rel. 40, Created)
 DT 20-AUG-2001 (Rel. 40, Last sequence update)
 DT 20-AUG-2001 (Rel. 40, Last annotation update)
 DE DUAL-SPECIFICITY TYROSINE-PHOSPHORYLATION REGULATED KINASE 2
 DE (EC 2.7.1.-).
 GN DYRK2.
 OS Homo sapiens (Human).

Query Match 78.0%; Score 32; DB 1; Length 528;
 Best Local Similarity 71.4%; Pred. No. 18;
 Matches 5; Conservative 1; Mismatches 1; Indels 0; Gaps 0;
 QY 1 KKRIMHC 7
 DB 268 KNRIHC 274
 RESULT 9
 ID_AMPI_LYCES STANDARD; PRT; 571 AA.
 AC Q10712; Q9S9A3;
 DT 01-NOV-1997 (Rel. 35, Created)
 DT 01-NOV-1997 (Rel. 35, Last sequence update)
 DT 20-AUG-2001 (Rel. 40, Last annotation update)
 DE CHLOROPLAST AMINOPEPTIDASE 1 PRECURSOR (EC 3.4.11.1) (LEUCINE
 DE AMINOPEPTIDASE) (LAP) (LEUCYL AMINOPEPTIDASE) (PROLINE AMINOPEPTIDASE)
 DE (EC 3.4.11.5) (PROLYL AMINOPEPTIDASE) (DR57).

GN LAPAL OR LAP OR LAP2.
 OS Lycopersicon esculentum (Tomato).
 OC Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
 OC Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots;
 OC Asteridae; euasterids I; Solanales; Solanaceae; Solanum.
 OX NCBI_TaxID=4081;
 RP [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN=CV. Peto 238R; TISSUE=Leaf;
 RX MEDLINE=96421572; PubMed=8824220;
 RA Gu Y.Q., Chao W.S., Walling L.L.;
 RT "Localization and post-translational processing of the wound-induced
 RT leucine aminopeptidase proteins of tomato.";
 RL J. Biol. Chem. 271:25880-25887(1996).
 RN [2]
 RN SEQUENCE OF 49-571 FROM N.A.
 RC STRAIN=CV. VF36; TISSUE=Pistil;
 RX MEDLINE=95375233; PubMed=7647301;
 RA Milligan S.B., Gasser C.S.;
 RT "Nature and regulation of pistil-expressed genes in tomato.";
 RL Plant Mol. Biol. 28:691-711(1995).
 RN [3]
 RN SEQUENCE OF 103-571 FROM N.A.
 RC STRAIN=CV. Peto 238R; TISSUE=Leaf;
 RX MEDLINE=94052201; PubMed=8234334;
 RA Pautot V., Holzer F.M., Reisch B., Walling L.L.;
 RT "Leucine aminopeptidase: an inducible component of the defense
 RT response in Lycopersicon esculentum (tomato).";
 RL Proc. Natl. Acad. Sci. U.S.A. 90:9906-9910(1993).
 CC -1- FUNCTION: PRESUMABLY INVOLVED IN THE PROCESSING AND REGULAR
 CC TURNOVER OF INTRACELLULAR PROTEINS.
 CC -1- CATALYTIC ACTIVITY: RELEASE OF AN N-TERMINAL AMINO ACID, XAA-|-
 CC XBB-, IN WHICH XAA IS PREFERABLY LEU, BUT MAY BE OTHER AMINO ACIDS
 CC INCLUDING PRO ALTHOUGH NOT ARG OR LYS, AND XBB MAY BE PRO.
 CC -1- COFACTOR: BINDS TWO ZINC IONS (BY SIMILARITY).
 CC -1- SUBCELLULAR LOCATION: CHLOROPLAST.
 CC -1- INDUCTION: BY WOUNDING.
 CC -1- SIMILARITY: BELONGS TO PEPTIDASE FAMILY M17; ALSO KNOWN AS THE
 CC CYTOSOL AMINOPEPTIDASE FAMILY.
 CC -----
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 CC -----
 CC EMBL: U50151; AAC49456.1; .
 CC EMBL: U50152; AAC49457.1; .
 CC EMBL: U20593; AAC80498.1; .
 CC HSP: P00727; IBPW.
 CC MEROPS: M17.002; .
 DR InterPro: IPR000819; Peptidase_M17.
 DR Pfam: PF00883; Peptidase_M17. 1.
 DR PRINTS: PR00481; LAMNOPTDASE.
 DR PROSITE: PS00631; CYTOSOLAP; 1.
 KW Transit peptide; Chloroplast; Aminopeptidase; Hydrolase; Zinc.
 FT TRANSIT 1 53
 FT CHAIN 54 571
 FT DOMAIN 169 174
 FT METAL 342 342
 FT METAL 347 347
 FT METAL 367 367
 FT METAL 427 427
 FT METAL 429 429
 FT ACT_SITE 354 354
 FT ACT_SITE 431 431
 FT VARIANT 358 358
 FT CONFLICT 271 271
 FT CONFLICT 315 315
 FT CONFLICT 515 515
 FT CONFLICT 571 AA; 60279 MW; C7A224837E73939D CRC64;
 SQ SEQUENCE

Query Match 78.0%; Score 32; DB 1; Length 571;
 Best Local Similarity 83.3%; Pred. No. 20;
 Matches 5; Conservative 1; Mismatches 0; Indels 0; Gaps 0;
 QY 2 KRIMHC 7
 Db 48 KRIVHC 53
 ID AMPL_SOLITU STANDARD; PRT: 573 AA.
 AC P31427;
 DT 01-JUL-1993 (Rel. 26, Created)
 DT 01-OCT-1996 (Rel. 34, Last sequence update)
 DT 01-OCT-1996 (Rel. 34, Last annotation update)
 DE CHLOROPLAST AMINOPEPTIDASE PRECURSOR (EC 3.4.11.1) (LEUCINE
 DE AMINOPEPTIDASE) (LAP) (LEUCYL AMINOPEPTIDASE) (PROLINE AMINOPEPTIDASE)
 DE (EC 3.4.11.5) (PROLYL AMINOPEPTIDASE).
 GN LAP.
 OS Solanum tuberosum (Potato).
 OC Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
 OC Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots;
 OC Asteridae; euasterids I; Solanales; Solanaceae; Solanum.
 OX NCBI_TaxID=4113;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN=CV. DESIREE;
 RX MEDLINE=94339796; PubMed=7765119;
 RA Herbers K., Prat S., Willmitzer L.;
 RT "Functional analysis of a leucine aminopeptidase from Solanum
 RT tuberosum L.";
 RL Planta 194:230-240(1994).
 RN [2]
 RP SEQUENCE OF 19-573 FROM N.A.
 RC STRAIN=CV. DESIREE; TISSUE=Leaf;
 RX MEDLINE=93005746; PubMed=1392612;
 RA Hilgmann T., Ebner M., Pena-Cortes H., Sanchez-Serrano J.J.,
 RA Willmitzer L., Prat S.;
 RT "General roles of abscisic and jasmonic acids in gene activation as a
 RT result of mechanical wounding.";
 RL Plant Cell 4:1157-1170(1992).
 CC -1- FUNCTION: PRESUMABLY INVOLVED IN THE PROCESSING AND REGULAR
 CC TURNOVER OF INTRACELLULAR PROTEINS.
 CC -1- CATALYTIC ACTIVITY: RELEASE OF AN N-TERMINAL AMINO ACID, XAA-|-
 CC XBB-, IN WHICH XAA IS PREFERABLY LEU, BUT MAY BE OTHER AMINO ACIDS
 CC INCLUDING PRO ALTHOUGH NOT ARG OR LYS, AND XBB MAY BE PRO.
 CC -1- COFACTOR: BINDS TWO ZINC IONS (BY SIMILARITY).
 CC -1- SUBUNIT: HOMOHETEROMER (PROBABLE).
 CC -1- SUBCELLULAR LOCATION: CHLOROPLAST (BY SIMILARITY).
 CC -1- TISSUE SPECIFICITY: IN TUBERS AND FLORAL BUDS OF UNTREATED PLANTS.
 CC AFTER ABA TREATMENT OR MECHANICAL WOUNDING IS MOSTLY ACCUMULATED
 CC IN LEAVES, TO A LESSER EXTENT IN STEMS, BUT NOT IN ROOTS.
 CC -1- INDUCTION: BY ABSICISIC ACID (ABA), JASMONIC ACID (JA) AND
 CC WOUNDING.
 CC -1- SIMILARITY: BELONGS TO PEPTIDASE FAMILY M17; ALSO KNOWN AS THE
 CC CYTOSOL AMINOPEPTIDASE FAMILY.
 CC -----
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 CC -----
 CC EMBL: X77015; CAA54314.1; .
 CC EMBL: X67845; CAA8038.1; .
 CC PIR: S24769; S24769.
 CC PIR: PQ0470; PQ0470.
 CC HSSP: P00727; ILAN.

```
DR MEROPS; M17.002;
DR InterPro: IPR000819; Peptidase_M17.
DR Pfam: PF00883; Peptidase_M17.1.
DR PRINTS: PR00481; LAMNOPPTDASE.
DR PROSITE: PS00631; CYTOSOL_AP.1.
KW Transit peptide; Chloroplast; Aminopeptidase; Hydrolase; Zinc.
FT CHAIN 1 53
FT CLUST 53
FT DOMAIN 169 174 POLY-ALA.
FT METAL 342 342 ZINC (2) (BY SIMILARITY).
FT METAL 347 347 ZINC (1 AND 2) (BY SIMILARITY).
FT METAL 367 367 ZINC (2) (BY SIMILARITY).
FT METAL 427 427 ZINC (1) (BY SIMILARITY).
FT METAL 429 429 ZINC (1 AND 2) (BY SIMILARITY).
FT METAL 429 429 ZINC (1 AND 2) (BY SIMILARITY).
FT ACT_SITE 354 354 POTENTIAL.
FT ACT_SITE 431 431 POTENTIAL.
SQ SEQUENCE 573 AA; 60122 MW; 3152145A4A7FB291 CRC64;

Query Match 78.0%; Score 32; DB 1; Length 573;
Best Local Similarity 83.3%; Pred. No. 20;
Matches 5; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

QY 2 KRIMHC 7
DB 48 KRIVHC 53

RESULT 11
RL44_TRYBB STANDARD; PRT; 105 AA.
AC P17843;
DT 01-AUG-1990 (Rel. 15, Created)
DT 20-AUG-2001 (Rel. 40, Last sequence update)
DT 20-AUG-2001 (Rel. 40, Last annotation update)
DE 60S RIBOSOMAL PROTEIN L44.
GN RPL44.
OS Trypanosoma brucei brucei.
OC Eukaryota; Euglenozoa; Kinetoplastida; Trypanosomatidae; Trypanosoma.
OX NCBI_TaxID=5702;
RN [1]
RC SEQUENCE FROM N.A.
RP STRAIN=ETRO 1125;
RX MEDLINE=90251460; PubMed=2339065;
RA Tebabi P., Halleux S., Pays E.;
RT "Nucleotide sequence of a full-length cDNA coding for the ribosomal L44 protein of Trypanosoma brucei."
RL Nucleic Acids Res. 18:2809-2809(1990).
CC -1- SUBCELLULAR LOCATION: CYTOPLASMIC.
CC -1- SIMILARITY: BELONGS TO THE L44E FAMILY OF RIBOSOMAL PROTEINS.
CC -----
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CC or send an email to license@isb-sib.ch).
CC -----
CC EMBL; X52122; CNA36367.1;
CC PIR; S10012; R6UT8A.
CC InterPro: IPR000552; Ribosomal_L44E.
CC Pfam; PF00935; Ribosomal_L44; 2.
CC ProDom; PD002841; Ribosomal_L44E; 1.
CC PROSITE; PS01172; RIBOSOMAL_L44E; 1.
KW Ribosomal protein.
FT INIT_MET 0 0 BY SIMILARITY.
SQ SEQUENCE 105 AA; 12322 MW; FA19423F109E7819 CRC64;

Query Match 75.6%; Score 31; DB 1; Length 105;
Best Local Similarity 71.4%; Pred. No. 5.9;
Matches 5; Conservative 1; Mismatches 1; Indels 0; Gaps 0;
```

```
QY 1 KKRIMHC 7
DB 5 KKKKMHK 11

RESULT 12
Y091_NPVOP STANDARD; PRT; 279 AA.
ID Y091_NPVOP
AC O10341;
DT 01-NOV-1997 (Rel. 35, Created)
DT 01-NOV-1997 (Rel. 35, Last sequence update)
DT 01-NOV-1997 (Rel. 35, Last annotation update)
DE HYPOTHETICAL 29.3 KDA PROTEIN (ORF92).
OS Orgyia pseudotsugata multicausid polyhedrosis virus (OpMPNV).
OC Viruses; dsDNA viruses, no RNA stage; Baculoviridae;
OC Nucleopolyhedrovirus.
OX NCBI_TaxID=164623;
RN [1]
RC SEQUENCE FROM N.A.
RP MEDLINE=97271300; PubMed=9126251;
RX Ahrens C.H., Russell R.R., Funk C.J., Evans J., Harwood S.,
RA Rohrmann G.F.;
RT "The sequence of the Orgyia pseudotsugata multicausid polyhedrosis virus genome."
RL Virology 229:381-393(1997).
CC -1- SIMILARITY: TO CORRESPONDING ORF IN ACMPNV.
CC -----
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CC -----
CC EMBL; J75930; AAC59091.1;
CC KW Hypothetical protein.
SQ SEQUENCE 279 AA; 29289 MW; 6FA4DA01009DBF0 CRC64;

Query Match 75.6%; Score 31; DB 1; Length 279;
Best Local Similarity 66.7%; Pred. No. 16;
Matches 4; Conservative 2; Mismatches 0; Indels 0; Gaps 0;

QY 2 KRIMHC 7
DB 233 QRVMHC 238

RESULT 13
CYCH_XENLA STANDARD; PRT; 323 AA.
ID CYCH_XENLA
AC P51947;
DT 01-OCT-1996 (Rel. 34, Created)
DT 01-OCT-1996 (Rel. 34, Last sequence update)
DT 20-AUG-2001 (Rel. 40, Last annotation update)
DE CYCLIN H (MO15-ASSOCIATED PROTEIN) (P36).
OS Xenopus laevis (African clawed frog).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Amphibia; Batrachia; Anura; Mesobatrachia; Pipidae;
OC Xenopodinae; Xenopus.
OX NCBI_TaxID=8355;
RN [1]
RC SEQUENCE FROM N.A.
RP Martinez A.-M.;
RA Submitted (JAN-1995) to the EMBL/GenBank/DBJ databases.
RN [2]
RP PARTIAL SEQUENCE.
RC TISSUE=Oocyte;
RX MEDLINE=95045408; PubMed=7957080;
RA Labbe J.-C., Martinez A.-M., Pesquet D., Capony J.-P., Darbon J.-M.,
RA Derancourt J., Devault A., Morin N., Cavadore J.-C., Doree M.;
```

RT "p40MO15 associates with a p36 subunit and requires both nuclear
 RT translocation and Thr176 phosphorylation to generate cdk-activating
 RT kinase activity in Xenopus oocytes.";
 RL EMBO J. 13:5155-5164(1994).
 CC -!- FUNCTION: SEEMS TO BE A REGULATORY COMPONENT OF CDK-ACTIVATING
 CC KINASE (CAK).
 CC -!- SUBUNIT: ASSOCIATES WITH CDK7 AND MAT1.
 CC -!- SUBCELLULAR LOCATION: NUCLEAR.
 CC -!- SIMILARITY: BELONGS TO THE CYCLIN FAMILY. CYCLIN C SUBFAMILY.
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 CC -----
 CC EMBL: U20505; AAR62236.1; ..
 CC HSP: F51946; IJRW.
 CC InterPro: IPR000553; Cyclin.
 CC SMART: SM00385; CYCLIN; 1.
 CC PROSITE: PS00292; CYCLINS; FALSE-NEG.
 CC Cyclin; Cell cycle; Cell division; Nuclear protein.
 CC CONFLICT 266 266 R -> Y (IN REF. 2; AA SEQUENCE).
 CC SEQUENCE 323 AA; 37600 MW; 14BCDCA000843DC8 CRC64;

Query Match 75.6%; Score 31; DB 1; Length 323;
 Best Local Similarity 71.4%; Pred. No. 18;
 Matches 5; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

QY 1 KKRIMHC 7
 DB 274 KRLDHC 280
 |||: ||

RESULT 14

SUN_HAEIN STANDARD; PRT; 451 AA.
 ID SUN_HAEIN
 AC P44788;
 DT 01-NOV-1995 (Rel. 32, Created)
 DT 01-NOV-1995 (Rel. 32, Last sequence update)
 DT 20-AUG-2001 (Rel. 40, Last annotation update)
 DE SUN PROTEIN (PMU PROTEIN).
 GN SUN OR PMU OR HI0624.
 OS Haemophilus influenzae.
 OC Bacteria; Proteobacteria; gamma subdivision; Pasteurellaceae;
 OC Haemophilus.
 OX NCBI_TaxID=727;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN=PD / KW20 / ATCC 51907;
 RX MEDLINE=95350630; PubMed=7542800;
 RA Fleischmann R.D., Adams M.D., White O., Clayton R.A., Kirkness E.F.,
 RA Kierlavage A.R., Bult C.J., Tomb J.-F., Dougherty B.A., Merrick J.M.,
 RA McElveney K., Sutton G., Fitzhugh W., Fields C.A., Gocayne J.D.,
 RA Scott J.D., Shirley R., Liu L.-I., Glodek A., Kelley J.M.,
 RA Weidman J.F., Phillips C.A., Spriggs T., Hedblom E., Cotton M.D.,
 RA Utterback T.R., Hanna M.C., Nguyen D.T., Saudek D.M., Brandon R.C.,
 RA Fine L.D., Fritchman J.L., Fuhrmann J.L., Geoghagen N.S.M.,
 RA Gnehm C.L., McDonald L.A., Small K.V., Fraser C.M., Smith H.O.,
 RA Venter J.C.;
 RT "Whole-genome random sequencing and assembly of Haemophilus
 RT influenzae Rd.";
 RL Science 269:496-512(1995).
 CC -!- SIMILARITY: BELONGS TO THE SUN (BACTERIAL) / NUCLEOLAR PROTEIN
 CC NOLI/NOF2 (EUKARYOTES) FAMILY.
 CC -----
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 CC -----

CC EMBL: U32745; AAC22284.1; ..
 CC TIGR: HI0624;
 CC InterPro: IPR001678; Noli_Nop2_Sun.
 CC InterPro: IPR000139; Nusb.
 CC DR InterPro: IPR000051; SAM_bind.
 CC Pfam: PF01189; Noli_Nop2_Sun; 1.
 CC Pfam: PF01029; Nusb; 1.
 CC DR PRODOM: PD005242; Nusb; 1.
 CC DR PROSITE: PS01153; NOLI_NOP2_SUN; 1.
 CC Complete proteome.
 KW PROSITE: PS01153; NOLI_NOP2_SUN; 1.
 SQ SEQUENCE 451 AA; 50597 MW; D91FAB88FFDE34B0 CRC64;

Query Match 75.6%; Score 31; DB 1; Length 451;
 Best Local Similarity 71.4%; Pred. No. 25;
 Matches 5; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

QY 1 KKRIMHC 7
 DB 88 KTRIVHC 94
 |||: ||

RESULT 15

CUL3_HUMAN STANDARD; PRT; 768 AA.
 ID CUL3_HUMAN
 AC Q13618; Q75415; Q9UB18; Q9UET7;
 DT 01-NOV-1997 (Rel. 35, Created)
 DT 20-AUG-2001 (Rel. 40, Last sequence update)
 DT 20-AUG-2001 (Rel. 40, Last annotation update)
 DE CULLIN HOMOLOG 3 (CUL-3).
 GN CUL3 OR KIAA0617.
 OS Homo sapiens (Human).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 OC Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
 OX NCBI_TaxID=9606;
 RN [1]
 RP SEQUENCE FROM N.A.
 RX MEDLINE=98406061; PubMed=9733711;
 RA Du M., Sansores-Garcia L., Zu Z., Wu K.K.;
 RT "Cloning and expression analysis of a novel salicylate suppressible
 RT gene, HS-CUL-3, a member of cullin/Cdc53 family.";
 RL J. Biol. Chem. 273:24289-24292(1998).
 RN [2]
 RP SEQUENCE FROM N.A.
 RC TISSUE=Brain;
 RX MEDLINE=98403880; PubMed=9734811;
 RA Ishikawa K.-I., Nagase T., Suyama M., Miyajima N., Tanaka A.,
 RA Kotani H., Nomura N., Ohara O.;
 RT "Prediction of the coding sequences of unidentified human genes. X.
 RT The complete sequences of 100 new cDNA clones from brain which can
 RT code for large proteins in vitro.";
 RL DNA Res. 5:169-176(1998).
 RN [3]
 RP SEQUENCE FROM N.A.
 RC TISSUE=Colon carcinoma;
 RX MEDLINE=98326596; PubMed=9663463;
 RA Michel J.J., Xiong Y.;
 RT "Human CUL-1, but not other cullin family members, selectively
 RT interacts with SKP1 to form a complex with SKP2 and cyclin A.";
 RL Cell Growth Differ. 9:435-449(1998).
 RN [4]
 RP SEQUENCE OF 192-768 FROM N.A.
 RX MEDLINE=96279828; PubMed=8681378;
 RA Kipreos E.T., Lander L.E., Wing J.P., He W.W., Hedgecock E.M.;
 RT "cul-1 is required for cell cycle exit in C. elegans and identifies a
 RT novel gene family.";
 RL Cell 85:829-839(1996).
 RN [5]
 RP SEQUENCE OF 426-768 FROM N.A.

RC TISSUE=Brain;
RA Yu W., Sarginson J., Gibbs R.A.;
RL Submitted (MAR-1998) to the EMBL/GenBank/DBJ databases.
CC -!- SIMILARITY: BELONGS TO THE CULLIN FAMILY.
CC -----
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CC -----
DR EMBL; AF064087; AAC36304.1; -;
DR EMBL; AB014517; BAA31592.1; -;
DR EMBL; AF062537; AAC36682.1; -;
DR EMBL; U58089; AAC50546.1; -;
DR EMBL; AF052147; AAC28621.1; -;
DR MIM; 603136; -;
DR InterPro; IPR001373; Cullin.
DR InterPro; IPR002119; Histone_H2A.
DR Pfam; PF00888; Cullin; 1.
DR ProDom; PD000565; Histone_H2A; 1.
DR SMART; SM00182; CULLIN; 1.
DR PROSITE; PS01256; CULLIN_1; 1.
DR PROSITE; PS50089; CULLIN_2; 1.
FT CONFLICT 13 179 D -> G (IN REF. 3).
FT CONFLICT 159 179 DLRQTLLDMIAERKGEYVD -> GSSTANSIGYDKRAE
FT CONFLICT 426 451 DVFEYKQHLARLLTNKSVSDSE -> MYLNVINNTW
FT CONFLICT 451 QGDFSOIKVFLMTLK (IN REF. 5).
FT SEQUENCE 768 AA; 88930 MW; ALA02022480BF099 CRC64;

Query Match 75.6%; Score 31; DB 1; Length 768;
Best Local Similarity 66.7%; Pred. No. 43;
Matches 4; Conservative 2; Mismatches 0; Indels 0; Gaps 0;
QY 2 KRIMHC 7
Db :|||||
246 ERVMHC 251

Search completed: February 12, 2002, 12:04:02
Job time: 798 sec

GenCore version 4.5
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OM protein - protein search, using sw model

Run on: February 12, 2002, 11:51:39 ; Search time 55.4 seconds
(without alignments)
9.625 Million cell updates/sec

Title: US-09-606-129A-18

Perfect score: 41

Sequence: 1 KKRIMHC 7

Scoring table: BLOSUM62
Gapop 10.0 , Gapext 0.5

Searched: 219241 seqs, 76174552 residues

Total number of hits satisfying chosen parameters: 219241

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database :

1: pir1:*

2: pir2:*

3: pir3:*

4: pir4:*

Pred. No. is the number of results predicted by chance to have a
score greater than or equal to the score of the result being printed,
and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	41	100.0	295	2 A42268	biliverdin reducta
2	41	100.0	533	2 T04537	hypothetical prote
3	38	92.7	296	2 G02066	biliverdin reducta
4	38	92.7	296	2 S62624	biliverdin reducta
5	36	87.8	512	2 H86206	hypothetical prote
6	34	82.9	401	2 F85015	probable phosphati
7	34	82.9	431	2 T01723	1-phosphatidylinos
8	33	80.5	189	2 A38117	hypothetical prote
9	33	80.5	508	2 T02486	hypothetical prote
10	33	80.5	925	2 S0490	hypothetical prote
11	32	78.0	107	2 B64845	hypothetical prote
12	32	78.0	107	2 D85651	hypothetical prote
13	32	78.0	140	2 I49668	binding protein -
14	32	78.0	142	2 JC1365	FK506/rapamycin-bi
15	32	78.0	328	2 B65086	hydrogenase (EC 1.
16	32	78.0	328	2 A85959	hydrogenase-2 smal
17	32	78.0	377	2 A82213	methylocitrate synt
18	32	78.0	497	2 C86463	hypothetical prote
19	32	78.0	516	2 T23827	hypothetical prote
20	32	78.0	566	2 T07850	leucyl aminopeptid
21	32	78.0	571	2 T07849	leucyl aminopeptid
22	32	78.0	571	2 T07047	leucyl aminopeptid
23	32	78.0	573	1 S41376	leucyl aminopeptid
24	31	75.6	106	1 R6UT6A	ribosomal protein
25	31	75.6	137	2 T46904	hypothetical prote
26	31	75.6	144	2 H41700	C6 protein - rabbi
27	31	75.6	152	2 T32784	hypothetical prote
28	31	75.6	279	2 T10361	hypothetical prote
29	31	75.6	321	2 T48373	hypothetical prote

30 31 75.6 383 2 S51651 cyclin delta-2 - A
31 31 75.6 451 1 F64155 hypothetical prote
32 31 75.6 476 2 T17330 hypothetical prote
33 31 75.6 504 2 JC4775 interferon-induced
34 31 75.6 504 2 A56534 interferon-induced
35 31 75.6 861 2 T41945 primase - human he
36 31 75.6 905 2 B71562 probable oxoglutar
37 31 75.6 905 2 H81714 2-oxoglutarate deh
38 31 75.6 1687 2 T30176 EGF repeat transme
39 31 75.6 1687 2 T30176 protein-tyrosine-p
40 30 73.2 108 2 B61180 hypothetical prote
41 30 73.2 269 2 C71255 conserved hypothet
42 30 73.2 370 2 D84464 hypothetical prote
43 30 73.2 457 2 T21344 hypothetical prote
44 30 73.2 508 2 T22440 hypothetical prote
45 30 73.2 522 2 B86453 CDS protein F9L11.

ALIGNMENTS

RESULT 1

A42268

biliverdin reductase (EC 1.3.1.24) - rat

C:Species: Rattus norvegicus (Norway rat)

C>Date: 04-Mar-1993 #sequence_revision 18-Nov-1994 #text_change 05-Nov-1999

C:Accession: A42268

R:Fakhrai, H.; Maines, M.D.

J. Biol. Chem. 267, 4023-4029, 1992

A:Title: Expression and characterization of a cDNA for rat kidney biliverdin reduct

A:Reference number: A42268; MUID:92156147

A:Accession: A42268

A>Status: preliminary; not compared with conceptual translation

A:Molecule type: nucleic acid; protein

A:Residues: 1-295 <FAK>

A:Cross-references: GB:M81681; NID:g203177; PIDN:AAA40830.1; PID:g203178

A:Experimental source: kidney

A:Note: sequence extracted from NCBI backbone (NCBIP:82800)

C:Keywords: liver; oxidoreductase

Query Match

Best Local Similarity 100.0%; Score 41; DB 2; Length 295;

Matches 7; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 KKRIMHC 7

Db 274 KKRIMHC 280

RESULT 2

T04537

hypothetical protein F28J12.70 - Arabidopsis thaliana

C:Species: Arabidopsis thaliana (mouse-ear cress)

C>Date: 23-Apr-1999 #sequence_revision 23-Apr-1999 #text_change 14-May-1999

C:Accession: T04537

R:Bevan, M.; Hilbert, H.; Braun, M.; Holzer, E.; Brandt, A.; Duesterhoeft, A.; Ban

submitted to the protein sequence database, February 1998

A:Reference number: 215377

A:Accession: T04537

A:Molecule type: DNA

A:Residues: 1-633 <BEV>

A:Cross-references: EMBL:AL021710

A:Experimental source: cultivar Columbia; BAC clone F28J12

C:Genetics:

A:Map position: 4

A:introns: 281/3; 303/3; 442/1; 614/3

A:Note: F28J12.70

Query Match

Best Local Similarity 100.0%; Score 41; DB 2; Length 633;

Matches 7; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 KKRIMHC 7
|||||
Db 591 KKRIMHC 597

RESULT 3

G02066
blliverrdin reductase (EC 1.3.1.24) - human
N:Alternate names: blliverrdin IX-alpha reductase
C:Species: Homo sapiens (man)
C:Date: 21-Dec-1996 #sequence_revision 06-Jun-1997 #text_change 17-Jul-1998
C:Accession: G02066
R:Komuro, A.; Tobe, T.; Nakano, Y.; Yamaguchi, T.; Tomita, M.
submitted to the EMBL Data Library, August 1995
A:Reference number: H00768
A:Accession: G02066
A:Status: preliminary; translated from GB/EMBL/DBJ
A:Molecule type: mRNA
A:Residues: 1-296 <KOW>
A:Cross-references: EMBL:U34877; NID:g1143231; PID:g1143232
C:Keywords: oxidoreductase

Query Match 92.7%; Score 38; DB 2; Length 296;
Best Local Similarity 85.7%; Pred. No. 1.5;
Matches 6; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

QY 1 KKRIMHC 7
|||||
Db 275 KKRILHC 281

RESULT 4

S62624
blliverrdin reductase (EC 1.3.1.24) - human
N:Alternate names: blliverrdin IX-alpha reductase
C:Species: Homo sapiens (man)
C:Date: 28-Oct-1996 #sequence_revision 09-May-1997 #text_change 21-Jul-2000
C:Accession: S62624; S62622; S29736
R:Maines, M.D.; Polevoda, B.V.; Huang, T.J.; McCoubrey Jr., W.K.
Eur. J. Biochem. 235, 373-381, 1996
A:Title: Human blliverrdin IX-alpha reductase is a zinc-metalloprotein. Characterization
A:Reference number: S62622; MUID:96202961
A:Accession: S62624
A:Molecule type: mRNA
A:Residues: 1-296 <MAI>
A:Cross-references: EMBL:X93086; NID:g1246748; PIDN:CAA63635.1; PID:g1246749
A:Accession: S62622
A:Molecule type: protein
A:Residues: 3-24, 'X', 26-27, 'X', 29-36; 48-74; 228-234; 235-248 <MAF>
R:Maines, M.D.; Trakshel, G.M.
Arch. Biochem. Biophys. 300, 320-326, 1993
A:Title: Purification and characterization of human blliverrdin reductase.
A:Reference number: S29736; MUID:93143333
A:Accession: S29736
A:Molecule type: protein
A:Residues: 3-24, 'X', 26-27, 'X', 29-36; 48-74; 228-234; 235-248 <MAW>
A:Note: the sequence of peptide 1 from page 323 seems not to belong to this protein
C:Genetics:
A:Gene: BVR
C:Keywords: oxidoreductase
F:3-296/Product: blliverrdin reductase IX-alpha #status experimental <MAT>

Query Match 92.7%; Score 38; DB 2; Length 296;
Best Local Similarity 85.7%; Pred. No. 1.5;
Matches 6; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

QY 1 KKRIMHC 7
|||||
Db 275 KKRILHC 281

RESULT 5

H86206
hypothetical protein [imported] - Arabidopsis thaliana
C:Species: Arabidopsis thaliana (mouse-ear cress)
C:Date: 02-Mar-2001 #sequence_revision 02-Mar-2001 #text_change 31-Mar-2001
C:Accession: H86206
R:Theologis, A.; Ecker, J.R.; Palm, C.J.; Federspiel, N.A.; Kaul, S.; White, O.; Al
Chin, C.W.; Chung, M.K.; Conn, L.; Conway, A.B.; Creasy, T.H.; Dewar
ansen, N.F.; Hughes, B.; Huizar, L.
Nature 408, 816-820, 2000
A:Authors: Hunter, J.L.; Jenkins, J.; Johnson-Hopson, C.; Khan, S.; Khaykin, E.; Ki
C.A.; Li, J.H.; Li, Y.; Lin, X.; Liu, Z.A.; Luros, J.S.; Mafti, R.; Marz
Rizzo, M.; Rooney, T.; Rowley, D.; Sakano, H.
A:Authors: Salzberg, S.L.; Schwartz, J.R.; Shinn, P.; Southwick, A.M.; Sun, H.; Tal
ker, M.; Wu, D.; Yu, G.; Fraser, C.M.; Venter, J.C.; Davis, R.W.
A:Title: Sequence and analysis of chromosome 1 of the plant Arabidopsis.
A:Reference number: A86141; MUID:21016719
A:Accession: H86206
A:Status: preliminary
A:Molecule type: DNA
A:Residues: 1-512 <STO>
A:Cross-references: GB:AE005172; NID:g8954028; PIDN:AAF82202.1; GSPDB:GN00141
C:Genetics:
A:Map position: 1

Query Match 87.8%; Score 36; DB 2; Length 512;
Best Local Similarity 71.4%; Pred. No. 6.6;
Matches 5; Conservative 2; Mismatches 0; Indels 0; Gaps 0;

QY 1 KKRIMHC 7
|||||
Db 236 KKRLLHC 242

RESULT 6

F85015
probable phosphatidylinositol kinase [imported] - Arabidopsis thaliana
C:Species: Arabidopsis thaliana (mouse-ear cress)
C:Date: 16-Feb-2001 #sequence_revision 16-Feb-2001 #text_change 16-Feb-2001
C:Accession: F85015
R:anonymous, The European Union Arabidopsis Genome Sequencing Consortium, The Cold
Nature 402, 769-777, 1999
A:Title: Sequence and analysis of chromosome 4 of the plant Arabidopsis thaliana.
A:Reference number: A85001; MUID:20083488
A:Accession: F85015
A:Status: preliminary
A:Molecule type: DNA
A:Residues: 1-401 <STO>
A:Cross-references: GB:NC_001268; NID:g7267616; PIDN:CAB80928.1; GSPDB:GN00140
C:Genetics:
A:Gene: AT4g01190
A:Map position: 4

Query Match 82.9%; Score 34; DB 2; Length 401;
Best Local Similarity 85.7%; Pred. No. 14;
Matches 6; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 KKRIMHC 7
|||||
Db 353 KKRILHC 359

RESULT 7

T01723
1-phosphatidylinositol-4-phosphate 5-kinase type II homolog - Arabidopsis thaliana
N:Alternate names: protein A_IG002N01.9
C:Species: Arabidopsis thaliana (mouse-ear cress)
C:Date: 19-Feb-1999 #sequence_revision 19-Feb-1999 #text_change 24-Mar-1999
C:Accession: T01723
R:Scheet, P.; Maggi, L.

submitted to the EMBL Data Library, June 1997
 A:Description: The sequence of A. thaliana IG002N01.
 A:Reference number: Z14407
 A:Accession: T01723
 A>Status: translated from GB/EMBL/DBDJ
 A:Molecule type: DNA
 A:Residues: 1-431 <SCH>
 A:Cross-references: EMBL:AF007269; NID:g2191126; PID:g2191143
 A:Experimental source: cultivar Columbia
 C:Genetics:
 A:Map position: 4
 A:Introns: 40/2; 94/3; 161/3; 224/2; 255/1; 271/1; 303/1; 339/2
 A:Note: A_IG002N01.9

Query Match 82.9%; Score 34; DB 2; Length 431;
 Best Local Similarity 85.7%; Pred. No. 15;
 Matches 6; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 KKRIMHC 7
 |||||
 Db 393 KKRIEHC 389

RESULT 8
 A38117
 hypothetical protein 1 - Anabaena sp. insertion sequence IS895
 C:Species: Anabaena sp.
 C>Date: 24-Jul-1992 #sequence_revision 24-Jul-1992 #text_change 15-Oct-1999
 C:Accession: A38117
 R:Alam, J.; Vrba, J.M.; Cai, Y.; Martin, J.A.; Weislo, L.J.; Curtis, S.E.
 J. Bacteriol. 173, 5778-5783, 1991
 A:Title: Characterization of the IS895 family of insertion sequences from the cyanobacterium Anabaena sp. strain PCC 7120
 A:Reference number: A38117; MUID:91358370
 A:Accession: A38117
 A>Status: preliminary
 A:Molecule type: DNA
 A:Residues: 1-189 <CALA>
 A:Cross-references: GB:M67475; NID:g142026; PIDN:AAA98138.1; PID:g142027
 A:Experimental source: strain PCC 7120
 C:Genetics:
 A:Mobile element: insertion sequence IS895

Query Match 80.5%; Score 33; DB 2; Length 189;
 Best Local Similarity 71.4%; Pred. No. 11;
 Matches 5; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

QY 1 KKRIMHC 7
 |||||
 Db 171 KKRLKHC 177

RESULT 9
 T02486
 hypothetical protein At2g29990 [imported] - Arabidopsis thaliana
 N:Alternate names: hypothetical protein F23F1.9
 C:Species: Arabidopsis thaliana (mouse-ear cress)
 C>Date: 05-Mar-1999 #sequence_revision 05-Mar-1999 #text_change 16-Feb-2001
 C:Accession: T02486; B84703
 R:Rounsley, S.D.; Lin, X.; Ketchum, K.A.; Crosby, M.L.; Brandon, R.C.; Sykes, S.M.; Kaul
 submitted to the EMBL Data Library, August 1998
 A:Description: Arabidopsis thaliana chromosome II BAC F23F1 genomic sequence.
 A:Reference number: Z14675
 A:Accession: T02486
 A>Status: translated from GB/EMBL/DBDJ
 A:Molecule type: DNA
 A:Residues: 1-508 <ROU>
 A:Cross-references: EMBL:AC004680; NID:g3420043; PID:g3420052
 A:Experimental source: cultivar Columbia
 R:Lin, X.; Kaul, S.; Rounsley, S.D.; Shea, T.P.; Benito, M.I.; Town, C.D.; Fujii, C.Y.;
 M.; Koo, H.; Moffat, K.S.; Cronin, L.A.; Shen, M.; VanAken, S.E.; Umayam, L.; Tallon, L.;
 euss, D.; Nierman, W.C.; White, O.; Eisen, J.A.; Salzberg, S.L.; Fraser, C.M.; Venter, J.

Nature 402, 761-768, 1999
 A:Title: Sequence and analysis of chromosome 2 of the plant Arabidopsis thaliana.
 A:Reference number: A84420; MUID:20083487
 A:Accession: B84703
 A>Status: preliminary
 A:Molecule type: DNA
 A:Residues: 1-508 <STO>
 A:Cross-references: GB:AE002093; NID:g3420052; PIDN:AAC31853.1; GSPDB:GN00139
 C:Genetics:
 A:Gene: At2g29990; F23F1.9
 A:Map position: 2
 A:Introns: 158/3; 230/1; 283/3; 305/3; 360/2; 398/3; 458/3
 C:Superfamily: NADH dehydrogenase

Query Match 80.5%; Score 33; DB 2; Length 508;
 Best Local Similarity 57.1%; Pred. No. 28;
 Matches 4; Conservative 3; Mismatches 0; Indels 0; Gaps 0;

QY 1 KKRIMHC 7
 |||||
 Db 236 KRLLHC 242

RESULT 10
 S50490
 hypothetical protein YER032w - yeast (Saccharomyces cerevisiae)
 C:Species: Saccharomyces cerevisiae
 C>Date: 28-May-1993 #sequence_revision 24-Feb-1995 #text_change 23-Mar-2001
 C:Accession: S50490
 R:Dietrich, F.S.
 submitted to the EMBL Data Library, December 1994
 A:Description: The sequence of S. cerevisiae cosmid 9537, 9581, 9495, 9867, and 18
 A:Reference number: S50433
 A:Accession: S50490
 A:Molecule type: DNA
 A:Residues: 1-925 <DI>
 A:Cross-references: EMBL:U18778; NID:g603592; PIDN:AAB64565.1; PID:g603624; MIPS:YI
 C:Genetics:
 A:Gene: SGD:YFR1
 A:Cross-references: SGD:S0000834; MIPS:YER032w
 A:Map position: 5R

Query Match 80.5%; Score 33; DB 2; Length 925;
 Best Local Similarity 71.4%; Pred. No. 48;
 Matches 5; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

QY 1 KKRIMHC 7
 |||||
 Db 841 KKRLSHC 847

RESULT 11
 B64845
 hypothetical protein bl028 - Escherichia coli
 C:Species: Escherichia coli
 C>Date: 12-Sep-1997 #sequence_revision 17-Sep-1997 #text_change 18-Aug-2000
 C:Accession: B64845
 R:Blattner, F.R.; Plunkett III, G.; Bloch, C.A.; Perna, N.T.; Burland, V.; Riley, B.
 A.; Rose, D.J.; Mau, B.; Shao, Y.
 Science 277, 1453-1462, 1997
 A:Title: The complete genome sequence of Escherichia coli K-12.
 A:Reference number: A64720; MUID:97426617
 A:Accession: B64845
 A>Status: nucleic acid sequence not shown; translation not shown
 A:Molecule type: DNA
 A:Residues: 1-107 <BLAT>
 A:Cross-references: GB:AE000205; GB:U00096; NID:g1787265; PIDN:AAC74113.1; PID:g178
 A:Experimental source: strain K-12, substrain MG1655
 C:Superfamily: Escherichia coli hypothetical protein bl028

```

Query Match      78.0%; Score 32; DB 2; Length 107;
Best Local Similarity 57.1%; Pred. No. 11;
Matches 4; Conservative 3; Mismatches 0; Indels 0; Gaps 0;

QY 1 KKRIMHC 7
    ::|||
Db 90 QQRVMHC 96

RESULT 12
D85651
hypothetical protein Z1557 [imported] - Escherichia coli (strain O157:H7)
C:Species: Escherichia coli
C:Date: 16-Feb-2001 #sequence_revision 16-Feb-2001 #text_change 31-Mar-2001
C:Accession: D85651
R:Perna, N.T.; Plunkett III, G.; Burland, V.; Mau, B.; Glasner, J.D.; Rose, D.J.; Mayhew,
Miller, L.; Grotbeck, E.J.; Davis, N.W.; Lim, A.; Dimalanta, E.; Potamousis, K.; Apodaca,
Nature 409, 529-533, 2001
A:Title: Genome sequence of enterohemorrhagic Escherichia coli O157:H7.
A:Reference number: 885480; MUID:21074935; PMID:11206551
A:Accession: D85651
A:Status: preliminary
A:Molecule type: DNA
A:Residues: 1-107 <STC>
A:Cross-references: GB:AE005174; NID:g12514431; PIDN:AAG55672.1; GSPDB:GN00145; UWGP:Z15
A:Experimental source: strain O157:H7, substrain EDL933
C:Genetics:
A:Gene: Z1557
C:Superfamily: Escherichia coli hypothetical protein bl028

Query Match      78.0%; Score 32; DB 2; Length 107;
Best Local Similarity 57.1%; Pred. No. 11;
Matches 4; Conservative 3; Mismatches 0; Indels 0; Gaps 0;

QY 1 KKRIMHC 7
    ::|||
Db 90 QQRVMHC 96

RESULT 13
I49668
banding protein - mouse
C:Species: Mus musculus (house mouse)
C:Date: 02-Jul-1996 #sequence_revision 02-Jul-1996 #text_change 16-Jul-1999
C:Accession: I49668
R:Hendrickson, B.A.; Zhang, W.; Craig, R.J.; Jin, Y.
Gene 134, 271-275, 1993
A:Title: Structural organization of the genes encoding human and murine FK506-binding pr
A:Reference number: I49668; MUID:94085790
A:Accession: I49668
A:Status: preliminary; translated from GB/EMBL/DBJ
A:Molecule type: DNA
A:Residues: 1-140 <RES>
A:Cross-references: GB:M77831; NID:g433782; PIDN:AAA37631.1; PID:g433783
C:Genetics:
A:Gene: Fkbp13
A:Introns: 55/3; 93/2; 109/1; 121/1
C:Superfamily: BKBP-type peptidylprolyl isomerase; BKBP-type peptidylprolyl isomerase ho
F:47-94/Domain: BKBP-type peptidylprolyl isomerase homology <PPI>

Query Match      78.0%; Score 32; DB 2; Length 140;
Best Local Similarity 71.4%; Pred. No. 14;
Matches 5; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

QY 1 KKRIMHC 7
    |||
Db 34 KKRVDHC 40

RESULT 14
JC1365

```

```

FK506/rapamycin-binding protein FKBP13 precursor - human
C:Species: Homo sapiens (man)
C:Date: 30-Sep-1993 #sequence_revision 30-Sep-1993 #text_change 07-Aug-1998
C:Accession: JC1365; A39602
R:Dilella, A.G.; Hawkins, A.; Craig, R.J.; Schreiber, S.L.; Griffin, C.A.
Biochem. Biophys. Res. Commun. 189, 819-823, 1992
A:Title: Chromosomal band assignments of the genes encoding human FKBP12 and FKBP13
A:Reference number: JC1365; MUID:93112052
A:Accession: JC1365
A:Molecule type: DNA
A:Residues: 1-142 <DIL>
R:Lin, Y.J.; Albers, M.W.; Lane, W.S.; Blier, B.E.; Schreiber, S.L.; Burakoff, S.J.
Proc. Natl. Acad. Sci. U.S.A. 88, 6677-6681, 1991
A:Title: Molecular cloning of a membrane-associated human FK506- and rapamycin-bind
A:Reference number: A39602; MUID:91319747
A:Accession: A39602
A:Molecule type: mRNA
A:Residues: 1-20, 'S'; 23-142 <JIN>
A:Cross-references: GB:M65128
C:Genetics:
A:Gene: GDB:FKBP2
A:Cross-references: GDB:133728; OMIM:186946
A:Map position: 11q13.1-11q13.3
A:Introns: 57/3; 95/2; 111/1; 123/1
C:Superfamily: BKBP-type peptidylprolyl isomerase; BKBP-type peptidylprolyl isomera
C:Keywords: immunoregulation
F:1-22/Domain: signal sequence #status predicted <SIG>
F:23-142/Product: FK506/rapamycin-binding protein FKBP13 #status predicted <MAT>
F:49-96/Domain: BKBP-type peptidylprolyl isomerase homology <PPI>

Query Match      78.0%; Score 32; DB 2; Length 142;
Best Local Similarity 71.4%; Pred. No. 14;
Matches 5; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

QY 1 KKRIMHC 7
    |||
Db 36 KKRVDHC 42

RESULT 15
B65086
hydrogenase (EC 1.18.99.1) 2 small chain - Escherichia coli
C:Species: Escherichia coli
C:Date: 12-Sep-1997 #sequence_revision 17-Sep-1997 #text_change 17-Mar-2000
C:Accession: B65086; A55516
R:Blattner, F.R.; Plunkett III, G.; Bloch, C.A.; Perna, N.T.; Burland, V.; Riley, M.
A.; Rose, D.J.; Mau, B.; Shao, Y.
Science 277, 1453-1462, 1997
A:Title: The complete genome sequence of Escherichia coli K-12.
A:Reference number: A64720; MUID:97426617
A:Accession: B65086
A:Status: nucleic acid sequence not shown; translation not shown
A:Molecule type: DNA
A:Residues: 1-328 <BLAT>
A:Cross-references: GB:AE000382; GB:U00096; NID:g2367182; PIDN:AAC76032.1; PID:g178
A:Experimental source: strain K-12, substrain MG1655
R:Menon, N.K.; Chateaux, C.F.; Deravatian, M.; Wendt, J.C.; Shannugam, K.T.; Peck
J. Bacteriol. 176, 4416-4423, 1994
A:Title: Cloning, sequencing, and mutational analysis of the hyb operon encoding Es
A:Reference number: A55516; MUID:94292472
A:Accession: A55516
A:Status: preliminary
A:Molecule type: DNA
A:Residues: 'V', 2-328 <MEN>
A:Cross-references: GB:U09177
C:Genetics:
A:Gene: hybA
C:Superfamily: unassigned ferredoxin 2[4Fe-4S]-related proteins; ferredoxin 2[4Fe-4
C:Keywords: oxidoreductase
F:105-163/Domain: ferredoxin 2[4Fe-4S] homology <FER>

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Query Match 78.0%; Score 32; DB 2; Length 328;
 Best Local Similarity 71.4%; Pred. No. 30;
 Matches 5; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

QY 1 KKRIMHC 7
 ||: |||
 Db 109 KQCMHC 115

Search completed: February 12, 2002, 11:51:40
 Job time: 301 sec

GenCore version 4.5
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OM protein - protein search, using sw model

Run On: February 12, 2002, 11:49:43 ; Search time 98.92 Seconds
(without alignments)
5.242 Million cell updates/sec

Title: US-09-606-129a-18
Perfect score: 41
Sequence: 1 KKRIMHC 7

Scoring table: BLOSUM62
Gapop 10.0 , Gapext 0.5

Searched: 522463 seqs, 74073290 residues

Total number of hits satisfying chosen parameters: 522463

Minimum DB seq length: 0
Maximum DB seq length: 2000000000
Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

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22: /SIDSB/gcgdata/geneseq/geneseq/AA2001.DAT.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	36	87.8	285	21	Arabidopsis thalia
2	34	82.9	94	21	Murine INGB1 isofo
3	34	82.9	279	21	Murine P37ING1 pol
4	33	80.5	508	21	Arabidopsis thalia
5	33	80.5	533	21	Arabidopsis thalia
6	33	80.5	564	21	Arabidopsis thalia
7	32	78.0	40	17	Bovine FKBP-13 imm
8	32	78.0	83	22	Human colon cancer
9	32	78.0	94	21	Human INGB1 isofo
10	32	78.0	99	13	Bovine RPKBP. Bos
11	32	78.0	104	21	Human secreted pro

12	32	78.0	128	20	AAV03245	Amino acid sequenc
13	32	78.0	137	22	AA87656	Bovine mammary tis
14	32	78.0	141	17	AA93551	Human FKBP-13 immu
15	32	78.0	141	21	AA83247	Human colon cancer
16	32	78.0	235	22	AA84700	Amino acid sequenc
17	32	78.0	279	21	AA97244	Human P37ING1. Ho
18	32	78.0	279	22	AAE0676	Tumour suppressor
19	32	78.0	279	22	AA84698	Amino acid sequenc
20	32	78.0	294	18	AAW1919	Tumour suppressor
21	32	78.0	294	19	AAW79675	Human P37ING1 poly
22	32	78.0	294	20	AAW03244	Amino acid sequenc
23	32	78.0	294	22	AA84697	Amino acid sequenc
24	32	78.0	328	22	AA89806	E. coli growth and
25	32	78.0	528	19	AA64559	Human protein kina
26	32	78.0	528	22	AAE02011	Human YAK1 (hYAK1)
27	32	78.0	549	21	AA88781	Amino acid sequenc
28	32	78.0	565	20	AAW96316	Acidic leucine ami
29	32	78.0	571	20	AAW96315	Acidic leucine ami
30	31	75.6	504	18	AAW36140	Bovine p58 protein
31	31	75.6	550	19	AAW71468	Cercospora nicotia
32	31	75.6	550	21	AA99884	Cercospora nicotia
33	31	75.6	882	22	AA63551	A human alpha-2 ma
34	31	75.6	887	20	AAW15344	Tumour suppressor
35	31	75.6	887	20	AAW28995	Tumour suppressor
36	31	75.6	899	22	AA63550	A human alpha-2 ma
37	31	75.6	912	22	AA63549	A human alpha-2 ma
38	31	75.6	1508	22	AA63548	A human alpha-2 ma
39	30	73.2	158	21	AA25386	Pinus radiata cell
40	29	70.7	386	21	AA621025	Arabidopsis thalia
41	29	70.7	386	21	AA63432	Arabidopsis thalia
42	29	70.7	391	21	AA843612	Human cancer assoc
43	29	70.7	396	21	AAW44277	Human nucleic acid
44	29	70.7	396	22	AAW38840	Human polypeptide
45	29	70.7	407	22	AAW40626	Human polypeptide

ALIGNMENTS

RESULT 1
AA828710
ID AAG28710 standard; Protein; 285 AA.
XX AAG28710;
AC AAG28710;
XX 17-OCT-2000 (first entry)
XX Arabidopsis thaliana protein fragment SEQ ID NO: 34031.
XX Protein identification; signal transduction pathway; metabolic pathway;
XX hybridisation assay; genetic mapping; gene expression control; promoter;
XX termination sequence.
XX Arabidopsis thaliana.
XX PN EPI033405-A2.
XX PD 06-SEP-2000.
XX PF 25-FEB-2000; 2000EP-0301439.
XX PR 25-FEB-1999; 99US-0121825.
XX PR 05-MAR-1999; 99US-0123180.
XX PR 09-MAR-1999; 99US-0123548.
XX PR 23-MAR-1999; 99US-0125788.
XX PR 25-MAR-1999; 99US-0126264.
XX PR 29-MAR-1999; 99US-0126785.
XX PR 01-APR-1999; 99US-0127462.
XX PR 06-APR-1999; 99US-0128234.
XX PR 08-APR-1999; 99US-0128714.
XX PR 16-APR-1999; 99US-0129845.
XX PR 19-APR-1999; 99US-0130077.
XX PR 21-APR-1999; 99US-0130449.

PR	23-APR-1999;	99US-0130510.	PR	20-JUL-1999;	99US-0144632.
PR	23-APR-1999;	99US-0130891.	PR	20-JUL-1999;	99US-0144684.
PR	28-APR-1999;	99US-0131449.	PR	21-JUL-1999;	99US-0144814.
PR	30-APR-1999;	99US-0132048.	PR	21-JUL-1999;	99US-0145086.
PR	30-APR-1999;	99US-0132407.	PR	21-JUL-1999;	99US-0145088.
PR	04-MAY-1999;	99US-0132484.	PR	22-JUL-1999;	99US-0145085.
PR	05-MAY-1999;	99US-0132485.	PR	22-JUL-1999;	99US-0145087.
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PR	07-MAY-1999;	99US-0132863.	PR	22-JUL-1999;	99US-0145145.
PR	11-MAY-1999;	99US-0134256.	PR	23-JUL-1999;	99US-0145218.
PR	14-MAY-1999;	99US-0134218.	PR	23-JUL-1999;	99US-0145224.
PR	14-MAY-1999;	99US-0134219.	PR	26-JUL-1999;	99US-0145276.
PR	14-MAY-1999;	99US-0134221.	PR	27-JUL-1999;	99US-0145913.
PR	14-MAY-1999;	99US-0134370.	PR	27-JUL-1999;	99US-0145918.
PR	18-MAY-1999;	99US-0134768.	PR	27-JUL-1999;	99US-0145919.
PR	19-MAY-1999;	99US-0134941.	PR	28-JUL-1999;	99US-0145951.
PR	20-MAY-1999;	99US-0135124.	PR	02-AUG-1999;	99US-0146386.
PR	21-MAY-1999;	99US-0135353.	PR	02-AUG-1999;	99US-0146388.
PR	24-MAY-1999;	99US-0135629.	PR	03-AUG-1999;	99US-0146389.
PR	25-MAY-1999;	99US-0136021.	PR	03-AUG-1999;	99US-0147038.
PR	27-MAY-1999;	99US-0136392.	PR	04-AUG-1999;	99US-0147204.
PR	28-MAY-1999;	99US-0136782.	PR	04-AUG-1999;	99US-0147302.
PR	01-JUN-1999;	99US-0137222.	PR	05-AUG-1999;	99US-0147192.
PR	03-JUN-1999;	99US-0137528.	PR	05-AUG-1999;	99US-0147260.
PR	04-JUN-1999;	99US-0137502.	PR	06-AUG-1999;	99US-0147303.
PR	07-JUN-1999;	99US-0137724.	PR	06-AUG-1999;	99US-0147416.
PR	08-JUN-1999;	99US-0138094.	PR	09-AUG-1999;	99US-0147493.
PR	10-JUN-1999;	99US-0138540.	PR	09-AUG-1999;	99US-0147935.
PR	10-JUN-1999;	99US-01386847.	PR	10-AUG-1999;	99US-0148171.
PR	14-JUN-1999;	99US-0139119.	PR	11-AUG-1999;	99US-0148319.
PR	16-JUN-1999;	99US-0139452.	PR	12-AUG-1999;	99US-0148341.
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PR	17-JUN-1999;	99US-0139492.	PR	13-AUG-1999;	99US-0148684.
PR	18-JUN-1999;	99US-0139454.	PR	16-AUG-1999;	99US-0149368.
PR	18-JUN-1999;	99US-0139455.	PR	17-AUG-1999;	99US-0149375.
PR	18-JUN-1999;	99US-0139456.	PR	18-AUG-1999;	99US-0149426.
PR	18-JUN-1999;	99US-0139457.	PR	20-AUG-1999;	99US-0149722.
PR	18-JUN-1999;	99US-0139458.	PR	20-AUG-1999;	99US-0149723.
PR	18-JUN-1999;	99US-0139459.	PR	20-AUG-1999;	99US-0149929.
PR	18-JUN-1999;	99US-0139460.	PR	23-AUG-1999;	99US-0149902.
PR	18-JUN-1999;	99US-0139461.	PR	23-AUG-1999;	99US-0149930.
PR	18-JUN-1999;	99US-0139462.	PR	25-AUG-1999;	99US-0150566.
PR	18-JUN-1999;	99US-0139463.	PR	26-AUG-1999;	99US-0150884.
PR	18-JUN-1999;	99US-0139750.	PR	27-AUG-1999;	99US-0151065.
PR	18-JUN-1999;	99US-0139763.	PR	27-AUG-1999;	99US-0151066.
PR	21-JUN-1999;	99US-0139817.	PR	27-AUG-1999;	99US-0151080.
PR	22-JUN-1999;	99US-0139899.	PR	30-AUG-1999;	99US-0151303.
PR	23-JUN-1999;	99US-0140353.	PR	31-AUG-1999;	99US-0151438.
PR	23-JUN-1999;	99US-0140354.	PR	01-SEP-1999;	99US-0151930.
PR	24-JUN-1999;	99US-0140692.	PR	07-SEP-1999;	99US-0152363.
PR	28-JUN-1999;	99US-0140823.	PR	10-SEP-1999;	99US-0153070.
PR	29-JUN-1999;	99US-0140991.	PR	13-SEP-1999;	99US-0153758.
PR	30-JUN-1999;	99US-0141287.	PR	15-SEP-1999;	

PR 14-OCT-1999; 99US-0159637.
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 PR 18-OCT-1999; 99US-0159584.
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 PR 21-OCT-1999; 99US-0160767.
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 PR 22-OCT-1999; 99US-0160989.
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 PR 28-OCT-1999; 99US-0161920.
 PR 28-OCT-1999; 99US-0161992.
 PR 28-OCT-1999; 99US-0161993.
 PR 29-OCT-1999; 99US-0162142.

Query Match 87.8%; Score 36; DB 21; Length 285;

Best Local Similarity 71.4%; Pred. No. 14; Mismatches 0; Indels 0; Gaps 0;

QY 1 KKRIMHC 7
 Db 13 kkrilh 19
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RESULT 2
 AAY97243
 ID AAY97243 standard; Protein; 94 AA.

XX AC AAY97243;
 XX DT 19-DEC-2000 (first entry)
 DE DE Murine INGB1 isoform N-terminal peptide.
 KW p53; tumour; cancer; detection; antibody; hybridisation; exon 1b;
 KW INGI; ing1; p33ING1; p37ING1; oncogene; gene therapy; diagnosis;
 KW proliferation disorder; transformation; transformatio; transformed cell; mouse.
 XX OS Mus musculus.
 XX PN WO200046370-A1.
 XX PD 10-AUG-2000.
 XX PF 04-FEB-2000; 2000WO-US02959.
 XX PR 04-FEB-1999; 99US-0118941.
 XX PA (UNII) UNIV ILLINOIS FOUND.
 XX PI Gudkov A, Zeremski M, Gurova KV, Grigorian IA;
 XX WPI; 2000-491278/43.

DR WPI; 2000-491278/43.

XX PT Detecting nucleic acid encoding exon 1b of ing1, useful for diagnosing
 XX PT and treating cancer, comprises contacting sample with isolated nucleic
 XX PT acid comprising sequence of exon 1b and detecting hybridized products
 XX PS Claim 6; Fig 7a; 134pp; English.

XX CC Mutations in or loss of the p53 gene occur in more than 50% of
 XX CC human tumours and tumour cell lines, but functional inactivation of
 XX CC the p53 pathway occurs in a much larger proportion of tumours. In
 XX CC many cases the mechanism of functional inactivation of the p53 gene
 XX CC remains unknown but p53 has been found to act in cooperation with

CC INGI. Functional cooperation between INGI and p53 suggested that
 CC INGI encoded a tumour suppressor protein that functioned within the
 CC p53 pathway. This data suggested a possible role for INGI in head
 CC and neck cancers and chromosomal location of the INGI placed it
 CC within a region that is frequently rearranged in head and neck
 CC cancers. Large scale analysis of tumours involving INGI has not
 CC revealed mutations in INGI nor significant variations in its
 CC expression suggesting that INGI was not a useful gene to study in
 CC cancer etiology. However, alternative initiation exons of the ing1
 CC gene, each having their own promoter have been discovered.
 CC Expression of one promoter (1a) produces a protein identical to
 CC INGI. Expression of a second promoter (1b) produces a protein having
 CC an identical C-terminal fragment to INGI but an additional 104
 CC N-terminal amino acids. The newly discovered protein has been
 CC designated p37ING1 (Wild type: p33ING1). p37ING1 has the
 CC characteristics of an oncogene. When overexpressed in cells (even
 CC those expressing wild type p53) p37ING1 is able to cause
 CC proliferation or transformation of those cells. Thus detecting a
 CC nucleic acid encoding exon 1b of ing1 by hybridisation with an
 CC isolated nucleic acid having the sequence of exon 1b of ing1
 CC or its antisense sequence can identify individuals expressing the
 CC oncogenic form of ing1. Novel peptide sequences taken from the 104
 CC N-terminal peptide of p37ING1 can also be used to raise antibodies
 CC that can also be used in detection methods for the p37ING1 variant.
 CC The polypeptides may be useful in gene therapy for treatment of cell
 CC proliferation disorders, especially cancers and for diagnosing and
 CC studying cancers.
 XX SQ Sequence 94 AA;

Query Match 82.9%; Score 34; DB 21; Length 94;

Best Local Similarity 57.1%; Pred. No. 12; Mismatches 0; Indels 0; Gaps 0;

QY 1 KKRIMHC 7
 Db 67 kkrvlhc 73
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RESULT 3

AAY97242
 ID AAY97242 standard; Protein; 279 AA.

XX AC AAY97242;
 XX DT 19-DEC-2000 (first entry)
 DE DE Murine P37ING1 polypeptide.
 KW p53; tumour; cancer; detection; antibody; hybridisation; exon 1b;
 KW INGI; ing1; p33ING1; p37ING1; oncogene; gene therapy; diagnosis;
 KW proliferation disorder; transformation; transformatio; transformed cell; mouse.

XX OS Mus musculus.

XX PN WO200046370-A1.

XX PD 10-AUG-2000.

XX PF 04-FEB-2000; 2000WO-US02959.

XX PR 04-FEB-1999; 99US-0118941.

XX PA (UNII) UNIV ILLINOIS FOUND.

XX PI Gudkov A, Zeremski M, Gurova KV, Grigorian IA;

XX WPI; 2000-491278/43.

XX DR N-PSDB; AAA53790.

XX PT Detecting nucleic acid encoding exon 1b of ing1, useful for diagnosing
 XX PT and treating cancer, comprises contacting sample with isolated nucleic

PT acid comprising sequence of exon 1b and detecting hybridized products
 XX Disclosure; Fig 12; 134pp; English.

XX Mutations in or loss of the p53 gene occur in more than 50% of
 CC human tumours and tumour cell lines, but functional inactivation of
 CC the p53 pathway occurs in a much larger proportion of tumours. In
 CC many cases the mechanism of functional inactivation of the p53 gene
 CC remains unknown but p53 has been found to act in cooperation with
 CC INGI. Functional cooperation between INGI and p53 suggested that
 CC INGI encoded a tumour suppressor protein that functioned within the
 CC p53 pathway. This data suggested a possible role for INGI in head
 CC and neck cancers and chromosomal location of the INGI placed it
 CC within a region that is frequently rearranged in head and neck
 CC cancers. Large scale analysis of tumours involving INGI has not
 CC revealed mutations in INGI nor significant variations in its
 CC expression suggesting that INGI was not a useful gene to study in
 CC cancer etiology. However, alternative initiation exons of the INGI
 CC gene, each having their own promoter have been discovered.
 CC Expression of one promoter (1a) produces a protein identical to
 CC INGI. Expression of a second promoter (1b) produces a protein having
 CC an identical C-terminal fragment to INGI but an additional 104
 CC N-terminal amino acids. The newly discovered protein has been
 CC designated p37ING1 (Wild type: p37ING1). p37ING1 has the
 CC characteristics of an oncogene. When overexpressed in cells (even
 CC those expressing wild type p53) p37ING1 is able to cause
 CC proliferation or transformation of those cells. Thus detecting a
 CC nucleic acid encoding exon 1b of INGI by hybridisation with an
 CC isolated nucleic acid having the sequence of exon 1b of INGI
 CC or its antisense sequence can identify individuals expressing the
 CC oncogenic form of INGI. Novel peptide sequences taken from the 104
 CC N-terminal peptide of p37ING1 can also be used to raise antibodies
 CC that can also be used in detection methods for the p37ING1 variant.
 CC The polypeptides may be useful in gene therapy for treatment of cell
 CC proliferation disorders, especially cancers and for diagnosing and
 CC studying cancers.

XX Sequence 279 AA;

Query Match 82.9%; Score 34; DB 21; Length 279;

Best Local Similarity 57.1%; Pred. No. 34;

Matches 4; Conservative 3; Mismatches 0; Indels 0; Gaps 0;

QY 1 KKRIMHC 7

Db 67 KRRVLC 73

RESULT 4

ID AAG48146

XX AAG48146 standard; Protein; 508 AA.

AC AAG48146;

XX 18-OCT-2000 (first entry)

DE Arabidopsis thaliana protein fragment SEQ ID NO: 60770.

XX Protein identification; signal transduction pathway; metabolic pathway;

KW hybridisation assay; genetic mapping; gene expression control; promoter;

KW termination sequence.

XX Arabidopsis thaliana.

OS Arabidopsis thaliana.

XX EP1033405-A2.

XX 06-SEP-2000.

PD 25-FEB-2000; 2000EP-0301439.

PF 25-FEB-1999; 99US-0121825.

XX 05-MAR-1999; 99US-0123180.

PR 09-MAR-1999; 99US-0123548.
 PR 23-MAR-1999; 99US-0125788.
 PR 29-MAR-1999; 99US-0126264.
 PR 01-APR-1999; 99US-0126785.
 PR 06-APR-1999; 99US-0127462.
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 PR 07-SEP-1999; 99US-0152363.
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 PR 28-OCT-1999; 99US-0161993.
 PR 29-OCT-1999; 99US-0162142.

Query Match 80.5%; Score 33; DB 21; Length 508;
 Best Local Similarity 57.1%; Pred. No. 95;
 Matches 4; Conservative 3; Mismatches 0; Indels 0; Gaps 0;

QY 1 KKRIMHC 7
 Db 236 krllhc 242
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RESULT 5
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 ID AAG48145 standard; Protein; 533 AA.
 XX
 AC AAG48145;
 XX
 XX 18-OCT-2000 (first entry)
 DT
 XX Arabidopsis thaliana protein fragment SEQ ID NO: 60769.
 DE
 XX Protein identification; signal transduction pathway; metabolic pathway;
 KW hybridisation assay; genetic mapping; gene expression control; promoter;
 KW termination sequence.
 XX
 OS Arabidopsis thaliana.
 XX
 PN EP1033405-A2.
 XX
 XX 06-SEP-2000.
 PD
 XX 25-FEB-2000; 2000EP-0301439.
 PF
 XX 25-FEB-1999; 99US-0121825.
 PR 05-MAR-1999; 99US-0123180.
 PR 09-MAR-1999; 99US-0123548.
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PR 31-AUG-1999; 99US-0151438.
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PR 28-OCT-1999; 99US-0161920.
PR 28-OCT-1999; 99US-0161992.
PR 28-OCT-1999; 99US-0161993.
PR 29-OCT-1999; 99US-0162142.

Query Match 80.5%; Score 33; DB 21; Length 564;
Best Local Similarity 57.1%; Pred. NO. 1e+02;
Matches 4; Conservative 3; Mismatches 0; Indels 0; Gaps 0;

Qy 1 KKRIMHC 7

Db 292 krllhc 298

RESULT 7

AAR93552

ID AAR93552 standard; Peptide; 40 AA.

AC AAR93552;

XX

DT 25-JUN-1996 (first entry)

XX

DE Bovine FKBP-13 immunophilin N-terminal peptide.

XX

KW FKBP-13; immunophilin; FK506; rapamycin; rheumatoid arthritis;
diabetes; organ transplant; graft versus host disease;
immunosuppressant.

XX

OS Bos taurus.

XX

PN US5498597-A.

XX

PD 12-MAR-1996.

XX

PF 17-JAN-1992; 92US-0822966.

XX

PR 17-JAN-1992; 92US-0822966.

XX

(DAND) DANA FARBER CANCER INST INC.

PA

PA (HARD) HARVARD COLLEGE.

XX

PI Bierer BE, Burakoff SJ, Schreiber SL;

XX

WPI; 1996-159713/16.

XX

Purified mammalian FKBP-13 polypeptide capable of binding FK506

useful for identifying and studying immunosuppressant drugs

XX

PS Disclosure; Column 8; 12pp; English.

XX

CC This sequence encoding the bovine FKBP-13 N-terminal sequence
corresponds to the N-terminal sequence of human FKBP-13. FKBP-13
may be used for identifying immunosuppressant drugs, and may be
used in combination with immunosuppressant drugs for therapeutic
purposes in the treatment of autoimmune diseases e.g. rheumatoid
arthritis and type-I diabetes, organ transplant and graft versus
host disease. The recombinant form of the protein could be
potentially smaller and therefore easier to introduce into cells
than intact FKBP-13.

XX SQ Sequence 40 AA; Query Match 78.0%; Score 32; DB 17; Length 40;
 Best Local Similarity 71.4%; Pred. No. 14;
 Matches 5; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

QY 1 KKRIMHC 7
 III: II
 Db 14 kkrvdhc 20

RESULT 8
 AAG76114
 ID AAG76114 standard; Protein; 83 AA.
 XX AC AAG76114;
 XX AC
 XX 03-SEP-2001 (first entry)
 XX Human colon cancer antigen protein SEQ ID NO:6878.
 DE Human; colon cancer; colon cancer antigen; diagnosis; detection;
 XX colorectal carcinoma; chromosome 11.
 KW Homo sapiens.
 XX WO200122920-A2.
 PN 05-APR-2001.
 XX 28-SEP-2000; 2000WO-US26524.
 XX 29-SEP-1999; 99US-0157137.
 PR 03-NOV-1999; 99US-0163280.
 XX (HUMA-) HUMAN GENOME SCI INC.
 PA Ruben SM, Barash SC, Birse CE, Rosen CA;
 PI WPI; 2001-235357/24.
 DR N-PSDB; AAH35519.
 XX Nucleic acids encoding 4277 human colon cancer-associated polypeptides,
 useful for preventing, diagnosing and/or treating colorectal cancers -
 Claim 11; Page 8320-8322; 9803pp; English.
 CC AAH32943 to AAH37195 and AAG7788 represent human colon
 CC cancer-associated nucleic acid molecules (N) and proteins (P), where
 CC the proteins are collectively known as colon cancer antigens. The colon
 CC cancer antigens have cytostatic activity and can be used in gene
 CC therapy and vaccine production. N and P may be used in the prevention,
 CC diagnosis and treatment of diseases associated with inappropriate P
 CC expression. For example, N and P may be used to treat disorders
 CC in a patient's genome that affect the activity of P by expressing
 CC inactive proteins or to supplement the patients' own production of P.
 CC Additionally, N may be used to produce the colon cancer-associated Ps,
 CC by inserting the nucleic acids into a host cell and culturing the cell
 CC to express the proteins. N and P can be used in the prevention, diagnosis
 CC and treatment of colorectal carcinomas and cancers. AAH37196 to AAH37204
 CC and AAB77789 represent sequences used in the exemplification of the
 CC present invention.
 CC N.B. Pages 666 to 682 and page 7053 of the sequence listing were
 CC missing at time of publication, meaning no sequences are present for
 CC SEQ ID NO:1027 to 1052, 7921 and 7922.
 XX Sequence 83 AA;
 SQ Query Match 78.0%; Score 32; DB 22; Length 83;

Best Local Similarity 71.4%; Pred. No. 27;
 Matches 5; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

QY 1 KKRIMHC 7
 III: II
 Db 48 kkrvdhc 54

RESULT 9
 AAY97245
 ID AAY97245 standard; Protein; 94 AA.
 XX AC AAY97245;
 XX 19-DEC-2000 (first entry)
 XX Human INGB1 isoform N-terminal peptide.
 DE p53; tumour; cancer; detection; antibody; hybridisation; exon 1b;
 XX INGI; INGI; p37ING1; p37ING1; oncogene; gene therapy; diagnosis;
 KW proliferation disorder; transformation; transformed cell; mouse.
 XX Homo sapiens.
 OS WO200046370-A1.
 PN 10-AUG-2000.
 XX 04-FEB-2000; 2000WO-US02959.
 PF 04-FEB-1999; 99US-0118941.
 PR (UNII) UNIV ILLINOIS FOUND.
 PA Gudkov A, Zeremski M, Gurova KV, Grigorian IA;
 PI WPI; 2000-491278/43.
 DR Detecting nucleic acid encoding exon 1b of INGI, useful for diagnosing
 PT and treating cancer, comprises contacting sample with isolated nucleic
 PT acid comprising sequence of exon 1b and detecting hybridized products
 XX Claim 26; Figure 7a; 134pp; English.
 CC Mutations in or loss of the p53 gene occur in more than 50% of
 CC human tumours and tumour cell lines, but functional inactivation of
 CC the p53 pathway occurs in a much larger proportion of tumours. In
 CC many cases the mechanism of functional inactivation of the p53 gene
 CC remains unknown but p53 has been found to act in cooperation with
 CC INGI. Functional cooperation between INGI and p53 suggested that
 CC INGI encoded a tumour suppressor protein that functioned within the
 CC p53 pathway. This data suggested a possible role for INGI in head
 CC and neck cancers and chromosomal location of the INGI placed it
 CC within a region that is frequently rearranged in head and neck
 CC cancers. Large scale analysis of tumours involving INGI has not
 CC revealed mutations in INGI nor significant variations in its
 CC expression suggesting that INGI was not a useful gene to study in
 CC cancer etiology. However, alternative initiation exons of the INGI
 CC gene, each having their own promoter have been discovered.
 CC Expression of one promoter (1a) produces a protein identical to
 CC INGI. Expression of a second promoter (1b) produces a protein having
 CC an identical C-terminal fragment to INGI but an additional 104
 CC N-terminal amino acids. The newly discovered protein has been
 CC designated p37ING1 (Wild type: p37ING1). p37ING1 has the
 CC characteristics of an oncogene. When overexpressed in cells (even
 CC those expressing wild type p53) p37ING1 is able to cause
 CC proliferation or transformation of those cells. Thus detecting a
 CC nucleic acid encoding exon 1b of INGI by hybridisation with an
 CC isolated nucleic acid having the sequence of exon 1b of INGI
 CC or its antisense sequence can identify individuals expressing the
 CC oncogenic form of INGI. Novel peptide sequences taken from the 104
 CC N-terminal peptide of p37ING1 can also be used to raise antibodies
 CC that can also be used in detection methods for the p37ING1 variant.

CC The polypeptides may be useful in gene therapy for treatment of cell
 CC proliferation disorders, especially cancers and for diagnosing and
 CC studying cancers.

XX
 SQ Sequence 94 AA;

Query Match 78.0%; Score 32; DB 21; Length 94;
 Best Local Similarity 57.1%; Pred. No. 30;
 Matches 4; Conservative 3; Mismatches 0; Indels 0; Gaps 0;

QY 1 KKRIMHC 7
 |||: ||
 Db 67 kkrvldhc 73

RESULT 10

AAR28979
 ID AAR28979 standard; protein; 99 AA.

XX
 AC AAR28979;

DT 24-MAR-1993 (first entry)
 XX

DE Bovine RFKBP.

XX Rapamycin; FK506; binding protein; RFKBP; prollyl isomerase;
 KW immunosuppressant; cyclosporin A; macrolide; bovine; thymus; BRFKBP;
 KW cis-trans prollyl isomerase activity; FKBP12.

XX Bos taurus.

XX WO9219745-A.

PN 12-NOV-1992.

XX 07-MAY-1992; 92WO-US03993.

PR 08-MAY-1991; 91US-0697113.

XX (VERT-) VERTEX PHARM INC.

PI Harding MW;

XX WPI; 1992-398871/48.

XX New prollyl isomerase and rapamycin FK506 binding protein - useful
 PT for screening potential immunosuppressive cpds.

XX Disclosure; Fig 1; 30pp; English.

XX This sequence corresponds to a fragment of a rapamycin FK506 binding
 CC protein (RFKBP). RFKBP is a prollyl isomerase structurally related to
 CC FK506 which does not bind the immunosuppressive cyclosporin A. RFKBP
 CC binds FK506 and rapamycin with quantitatively significant selectivity.
 CC RFKBP may be used in screening assays to detect new immunosuppressants
 CC and to differentiate rapamycin-like cpds. from FK506-like cpds.

CC Rapamycin is a macrolide which is structurally related to FK506.
 CC This RFKBP has been isolated from bovine thymus (BRFKBP) and was found
 CC to be of low molecular weight, approx. 16,000, and to have cis-trans
 CC prollyl isomerase activity. The N terminal of BRFKBP has been shown to
 CC have over 50% homology to the N terminal of FKBP12.

XX Sequence 99 AA;

Query Match 78.0%; Score 32; DB 13; Length 99;
 Best Local Similarity 71.4%; Pred. No. 32;
 Matches 5; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

QY 1 KKRIMHC 7
 |||: ||
 Db 14 kkrvldhc 20

RESULT 11

AAG03758
 ID AAG03758 standard; Protein; 104 AA.

XX
 AC AAG03758;

DT 06-OCT-2000 (first entry)
 XX

DE Human secreted protein, SEQ ID NO: 7839.

XX Human; 5' EST; expressed sequence tag; secreted protein; cDNA isolation;
 KW gene therapy; chromosome mapping.

XX Homo sapiens.

XX EPI033401-A2.

PN 06-SEP-2000.

PD 21-FEB-2000; 2000EP-0200610.

XX 26-FEB-1999; 99US-0122487.

PR (BEST) GENSET.

XX Dumas Milne Edwards J, Duclert A, Giordano J;

XX WPI; 2000-500381/45.

DR N-PSDB; AAC03764.

XX New nucleic acid that is a 5' expressed sequence tag (5' EST) for
 PT obtaining cDNAs and genomic DNAs that correspond to 5' ESTs and for
 PT diagnostic, forensic, gene therapy and chromosome mapping procedures -
 XX Claim 13; SEQ ID 7839; 71pp + CD-ROM; English.

XX The present sequence is a polypeptide encoded by one of a large number
 CC of 5' ESTs derived from mRNAs encoding secreted proteins. The 5' ESTs
 CC were prepared from total human RNAs or polyA+ RNAs derived from 30
 CC different tissues. EST sequences usually correspond mainly to the 3'
 CC untranslated region (UTR) of the mRNA because they are often obtained
 CC from oligo-dT primed cDNA libraries. Such ESTs are not well suited for
 CC isolating cDNA sequences derived from the 5' ends of mRNAs and even in
 CC those cases where longer cDNA sequences have been obtained, the full 5'
 CC UTR is rarely included. 5' ESTs are derived from mRNAs with intact 5'
 CC ends and can therefore be used to obtain full length cDNAs and genomic
 CC DNAs. 5' ESTs are also used in diagnostic, forensic, gene therapy and
 CC chromosome mapping procedures. They are used to obtain upstream
 CC regulatory sequences and to design expression and secretion vectors.

XX Sequence 104 AA;

Query Match 78.0%; Score 32; DB 21; Length 104;
 Best Local Similarity 71.4%; Pred. No. 33;
 Matches 5; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

QY 1 KKRIMHC 7
 |||: ||
 Db 36 kkrvldhc 42

RESULT 12

AAV03245
 ID AAV03245 standard; Protein; 128 AA.

XX
 AC AAV03245;

XX 26-AUG-1999 (first entry)
 DT

DE Amino acid sequence of the human p28-ING1 protein.

XX Human; p33-ING1 protein; growth regulation; apoptosis; DNA damage;
 KW inhibition; anchorage independent growth; cytotoxic drug; cancer;
 KW transcriptional activation; immortal cell line; p28-ING1 protein.
 XX
 OS Homo sapiens.
 XX
 FH Key Location/Qualifiers
 FT Misc_Difference 59..259
 FT /note= "p26-ING1 fragment"
 XX
 PN WO9916790-A1.
 XX
 XX 08-APR-1999.
 XX
 XX 24-SEP-1998; 98WO-US18179.
 XX
 XX 14-JAN-1998; 98US-0006783.
 PR 26-SEP-1997; 97US-0060138.
 XX
 XX (UNII) UNIV ILLINOIS BOARD OF TRUSTEES.
 PA (UYTE-) UNIV TECHNOLOGIES INT INC.
 XX
 PI Garkavtsev I, Gudkov A, Riabowol K;
 XX WPI; 1999-263685/22.
 XX
 XX Use of p33-ING1 peptides
 PT
 PS Example 8; Page 61; 64pp; English.
 XX
 XX This is the amino acid sequence of the human p28-ING1 protein,
 CC used in the method of the invention, involving the human p33-ING1
 CC protein. The ING1 gene encodes p33-ING1 which can be used to
 CC modulate the activity of, isolate or detect p53. Expression of the
 CC ING1 and p53 genes in a mammalian cell results in normal growth
 CC regulation anchorage-dependent growth and apoptosis as a response
 CC to irreversible DNA damage and other cellular insult. Inhibition of
 CC expression of either gene results in a loss of cellular growth
 CC control, anchorage independent growth, inhibition of apoptosis
 CC and resistance to radiation and cytotoxic drugs. The p33-ING1 is a
 CC component of the p53 signalling pathway that cooperates with p53 in
 CC negative regulation of cell proliferation by modulating p53 dependent
 CC transcriptional activation. Biological function of p53 signalling
 CC pathway can therefore be regulated (both enhanced or suppressed) by
 CC modulating p33-ING1 activity. The modulation of p33-ING1 activity can
 CC be used for the stimulation or restoration of the p53 pathway in
 CC anti cancer therapy or for the suppression of the p53 pathway to
 CC defend sensitive tissues from genotoxic stress or for the generation
 CC of immortal cell lines.
 XX
 XX Sequence 128 AA;
 SQ
 Query Match 78.0%; Score 32; DB 20; Length 128;
 Best Local Similarity 57.1%; Pred. No. 41;
 Matches 4; Conservative 3; Mismatches 0; Indels 0; Gaps 0;
 QY 1 KKRIMHC 7
 Db 67 kkrmlhc 73
 RESULT 13
 AAB87656
 ID AAB87656 standard; protein; 137 AA.
 XX
 AC AAB87656;
 XX
 DT 15-MAY-2001 (first entry)
 XX
 DE Bovine mammary tissue derived protein #47.
 XX

KW Bovine; mammary gland; cancer; tumour; angiogenesis.
 XX
 OS Bos taurus.
 XX
 PN WO200114553-A1.
 XX
 XX 01-MAR-2001.
 XX
 XX 23-AUG-2000; 2000WO-NZ00166.
 PF
 XX 23-AUG-1999; 99US-0150330.
 PR
 XX (GENE-) GENESIS RES & DEV CORP LTD.
 PA (NZPA-) NEW ZEALAND PASTORAL AGRIC RES INST LTD.
 XX
 PI Havukkala IJ, Gleen M, Grigor MR, Molenaar AJ;
 XX WPI; 2001-226619/23.
 DR
 XX New polypeptides and polynucleotides encoding the polypeptides, which
 PT are expressed in bovine mammary gland tissue, useful for stimulating
 PT mammary gland growth or function, or inducing differentiation of milk
 PT producing cells
 XX
 PS Claim 11; Page 80; 97pp; English.
 XX
 CC The present invention relates to proteins derived from bovine
 CC mammary gland cells. The invention is useful for stimulating the
 CC bovine mammary gland cell growth and function, inhibiting the
 CC growth of various mammary gland cancer cells, inhibiting
 CC angiogenesis and vascularization of tumours, or modulating
 CC the growth of blood vessels in a mammal.
 XX
 SQ Sequence 137 AA;
 Query Match 78.0%; Score 32; DB 22; Length 137;
 Best Local Similarity 71.4%; Pred. No. 43;
 Matches 5; Conservative 1; Mismatches 1; Indels 0; Gaps 0;
 QY 1 KKRIMHC 7
 Db 43 kkrvdhc 49
 RESULT 14
 AAR93551
 ID AAR93551 standard; Protein; 141 AA.
 XX
 AC AAR93551;
 XX
 DT 25-JUN-1996 (first entry)
 XX
 DE Human FKBP-13 immunophilin protein.
 XX
 KW FKBP-13; immunophilin; FK506; rapamycin; rheumatoid arthritis;
 KW diabetes; organ transplant; graft versus host disease;
 KW immunosuppressant.
 XX
 OS Homo sapiens.
 XX
 FH Key Location/Qualifiers
 FT Peptide 1..21
 FT /label= sig_peptide
 FT Misc-difference 22..61
 FT /note= "corresponds to bovine thymus FKBP-13 N-
 FT terminal sequence"
 FT Misc-difference 117..120
 FT /note= "endoplasmic reticulum retention sequence"
 XX
 PN US5498597-A.
 XX
 XX 12-MAR-1996.
 PD

```

XX 17-JAN-1992; 92US-0822966.
XX
XX 17-JAN-1992; 92US-0822966.
XX
XX (DAND ) DANA FARBER CANCER INST INC.
XX (HARD ) HARVARD COLLEGE.
XX
XX Brierer BE, Burakoff SJ, Schreiber SL;
XX WPI; 1996-159713/16.
XX DR N-PSDB; AAT18037.
XX
XX Purified mammalian FKBP-13 polypeptide capable of binding FK506
XX useful for identifying and studying immunosuppressant drugs
XX
XX Claim 1; Fig.1; 12pp; English.
XX
XX The FKBP-13 protein may be used for identifying immunosuppressant
XX drugs, and may be used in combination with immunosuppressant drugs
XX for therapeutic purposes in the treatment of autoimmune diseases e.g.
XX rheumatoid arthritis and type-1 diabetes, organ transplant and
XX graft versus host disease. The recombinant form of the protein
XX could be potentially smaller and therefore easier to introduce
XX into cells than intact FKBP-13.
XX
XX Sequence 141 AA;
XX
Query Match 78.0%; Score 32; DB 17; Length 141;
Best Local Similarity 71.4%; Pred. NO. 45;
Matches 5; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

QY 1 KKRIMHC 7
Db 35 kkrvdhc 41

RESULT 15
AAB53247
ID AAB53247 standard; Protein; 141 AA.
XX
XX AAB53247;
XX
XX 09-MAR-2001 (first entry)
XX
XX Human colon cancer antigen protein sequence SEQ ID NO:787.
XX
XX Human; colon cancer; colon cancer antigen; diagnosis; detection;
XX identification; cytostatic; cardioactive; neuroprotective; vulnerary;
XX immunomodulatory; muscular; gynaecological; gastrointestinal;
XX nephrotropic; antiinfective; antibacterial; gene therapy; wound;
XX neural disorder; immune system disorder; muscular disorder;
XX reproductive disorder; gastrointestinal disorder; renal disorder;
XX infectious disease; cardiovascular disorder.
XX
XX Homo sapiens.
XX
XX WO200055351-A1.
XX
XX 21-SEP-2000.
XX
XX 08-MAR-2000; 2000WO-US05883.
XX
XX 12-MAR-1999; 99US-0124270.
XX
XX (HUMA-) HUMAN GENOME SCI INC.
XX
XX Rosen CA, Ruben SM;
XX
XX WPI; 2000-587534/55.
XX DR N-PSDB; AAC98004.
XX

```

```

PT Colon cancer associated gene sequences, referred to as colon cancer
PT antigens, useful for the treatment, prevention, and diagnosis of colon
PT disorders such as colon cancer -
XX
XX Claim 11; Page 1346; 2104pp; English.
XX
XX AAC97991 to AAC98763 encode the human colon cancer associated proteins,
XX called human colon cancer antigens, given in AAB53234 to AAB54006. The
XX human colon cancer antigens can have cytostatic, cardioactive, muscular;
XX neuroprotective, immunomodulatory, gynaecological, gastrointestinal,
XX vulnerary, nephrotropic, antiinfective and antibacterial activities, and
XX can be used in gene therapy. The colon cancer antigen polynucleotides,
XX proteins and antibodies to the proteins are useful for the prevention,
XX treatment and diagnosis of colon disorders, such as colon cancer. The
XX polynucleotides may be used in diagnostics and research, such as for
XX chromosome identification, and as hybridisation probes. The proteins
XX may also be used to prevent diseases such as neural disorders, immune
XX system disorders, muscular disorders, reproductive disorders,
XX gastrointestinal disorders, wounds, renal disorders, infectious
XX diseases, and cardiovascular disorders. AAC98764 to AAC98772 and
XX AAB54007 represent sequences used in the exemplification of the present
XX invention.
XX
XX Sequence 141 AA;
XX

```

```

Query Match 78.0%; Score 32; DB 21; Length 141;
Best Local Similarity 71.4%; Pred. NO. 45;
Matches 5; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

QY 1 KKRIMHC 7
Db 27 knriihc 33

```

Search completed: February 12, 2002, 11:49:43
Job time: 499 sec

GenCore version 4.5
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OM protein - protein search, using sw model

Run on: February 12, 2002, 12:03:25 ; Search time 94.82 Seconds
(without alignments)
12.341 Million cell updates/sec

Title: US-09-606-129A-19
Perfect score: 46
Sequence: 1 QKLCHQKK 8

Scoring table: BLOSUM62
Gapop 10.0 , Gapext 0.5

Searched: 473505 seqs, 146273229 residues 473505
Total number of hits satisfying chosen parameters:

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database :

SPTREMBL_17:*
1: sp.archaea:*
2: sp.bacteria:*
3: sp.fungi:*
4: sp.human:*
5: sp.invertebrate:*
6: sp.mammal:*
7: sp.mhc:*
8: sp.organelle:*
9: sp.phage:*
10: sp.plant:*
11: sp.rodent:*
12: sp.virus:*
13: sp.vertebrate:*
14: sp.unclassified:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	37	80.4	636	12 090142	090142 spodoptera
2	37	80.4	1063	12 09J844	Q9J844 spodoptera
3	35	76.1	295	11 09CY64	Q9CY64 mus musculus
4	35	76.1	303	11 09PD21	Q9PD21 mus musculus
5	35	76.1	420	10 022019	022019 cyanidiosch
6	34	73.9	288	5 094164	Q94164 caenorhabdi
7	34	73.9	322	4 09BSM2	Q9BSM2 homo sapien
8	34	73.9	409	11 09D3R6	Q9D3R6 mus musculus
9	34	73.9	522	4 09H8F4	Q9H8F4 homo sapien
10	34	73.9	547	3 074308	074308 schizosacch
11	34	73.9	790	4 09NVK9	Q9NVK9 homo sapien
12	34	73.9	1188	5 096143	096143 plasmodium
13	34	73.9	1293	3 013348	013348 magnaporthe
14	34	73.9	1473	11 063625	063625 rattus norv
15	34	73.9	1654	4 09PIY6	Q9PIY6 homo sapien
16	33	71.7	261	4 09HIN6	Q9HIN6 homo sapien
17	33	71.7	263	1 09YDU2	Q9YDU2 aeropyrum p
18	33	71.7	266	2 096301	096301 alcaligenes
19	33	71.7	273	12 09PYW7	Q9PYW7 xestia c-ni

20	33	71.7	445	10 092PL5	Q92PL5 nicotiana t
21	33	71.7	540	8 09MT39	Q9MT39 solanum tub
22	33	71.7	566	4 09H2S0	Q9H2S0 homo sapien
23	33	71.7	588	4 09ULU1	Q9ULU1 homo sapien
24	33	71.7	673	11 09QVT4	Q9QVT4 mus sp. mpl
25	33	71.7	773	10 080631	080631 arabidopsis
26	33	71.7	1247	5 017461	Q17461 caenorhabdi
27	32	69.6	87	5 09NSN6	Q9NSN6 caenorhabdi
28	32	69.6	163	2 09EUG2	Q9EUG2 escherichia
29	32	69.6	312	11 09CSE2	Q9CSE2 mus musculu
30	32	69.6	336	4 09H495	Q9H495 homo sapien
31	32	69.6	380	5 021866	Q21866 caenorhabdi
32	32	69.6	397	4 09H497	Q9H497 homo sapien
33	32	69.6	397	4 09H6E7	Q9H6E7 homo sapien
34	32	69.6	422	10 09SSP1	Q9SSP1 arabidopsis
35	32	69.6	462	5 09UOE3	Q9UOE3 strongyloce
36	32	69.6	465	5 09XW94	Q9XW94 caenorhabdi
37	32	69.6	474	10 080588	080588 arabidopsis
38	32	69.6	548	5 020367	Q20367 caenorhabdi
39	32	69.6	590	2 045490	Q45490 bacillus st
40	32	69.6	628	12 090139	Q90139 mamestra br
41	32	69.6	828	11 09Q207	Q9Q207 mus musculu
42	32	69.6	828	11 09D6I6	Q9D6I6 mus musculu
43	32	69.6	843	12 0994E8	Q994E8 porcine ade
44	32	69.6	864	10 09C987	Q9C987 arabidopsis
45	32	69.6	920	11 09JJK7	Q9JJK7 mesocricetu

ALIGNMENTS

RESULT 1
090142 ID 090142 PRELIMINARY; PRT; 636 AA.
AC 090142;
DT 01-NOV-1998 (Tremblrel. 08, Created)
DT 01-NOV-1998 (Tremblrel. 08, Last sequence update)
DT 01-JUN-2001 (Tremblrel. 17, Last annotation update)
DE DNA POLYMERASE (FRAGMENT).
GN DPOL.
OS Spodoptera exigua nucleopolyhedrovirus.
OC Viruses; dsDNA viruses, no RNA stage; Baculoviridae;
OC Nucleopolyhedrovirus.
OX NCBI_TaxID=10454;
RN [1]
RP SEQUENCE FROM N.A.
RA Bulach D.M., Kumar C.A., Zaia A., Liang B., Tribe D.E.;
RT "Group II Nucleopolyhedrovirus Subgroups Revealed by Phylogenetic
RT Analysis of Polyhedrin and DNA Polymerase Gene Sequences."
RL Submitted (MAY-1998) to the EMBL/GenBank/DBJ databases.
CC -!- CATALYTIC ACTIVITY: N DEOXYNUCLEOSIDE TRIPHOSPHATE = N
CC PYROPHOSPHATE + DNA(N).
CC -!- SIMILARITY: BELONGS TO DNA POLYMERASE TYPE-B FAMILY.
DR EMBL; AF068186; AAC3749.1; -
DR InterPro; IPR002064; DNA_pol_B.
DR Pfam; PF00136; DNA_pol_B; 2.
DR PRINTS; PR00106; DNAPOLB.
DR SMART; SM00486; POLBC; 1.
KW DNA replication; DNA-binding; DNA-directed DNA polymerase.
FT NON_TER 1
FT NON_TER 636
SQ SEQUENCE 636 AA; 73836 MW; 9BDB36475B07DCD1 CRC64;

Query Match 80.4%; Score 37; DB 12; Length 636;
Best Local Similarity 85.7%; Pred. No. 12;
Matches 6; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

QY 2 KLCHQKK 8

DB 392 KLCHQKK 398

QY 1 QKLCQKK 8
|:|||||:
Db 288 QRLCHRKQ 295

RESULT 5
Q22019 PRELIMINARY; PRT; 420 AA.
AC Q22019;
DT 01-JAN-1998 (T-EMBLrel. 05, Created)
DT 01-JAN-1998 (T-EMBLrel. 05, Last sequence update)
DT 01-JUN-2001 (T-EMBLrel. 17, Last annotation update)
DE OSF420.
OS Cyanidioschyzon merolae.
OC Eukaryota; Rhodophyta; Bangiophyceae; Cyanidioschyzon.
OX NCBI_TaxID=45157;
RN [1]
RP SEQUENCE FROM N.A.
RA Ohta N.;
RL J. Plant Res. 110:235-245(1997).
DR EMBL; D63675; BAA22815.1;
DR Mendel; 23995; Cyame; 3036; 23995.
DR InterPro; IPR000178; IF2
DR InterPro; IPR001950; SUI1.
DR Pfam; PF02131; IF2; 1.
DR ProDom; PD186100; IF2; 1.
DR PROSITE; PS01118; SUI1.1; UNKNOWN.1.
SQ SEQUENCE 420 AA; 47691 MW; A6CAE107B24B4E19 CRC64;

Query Match 76.1%; Score 35; DB 10; Length 420;
Best Local Similarity 85.7%; Pred. No. 23;
Matches 6; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 QKLCQKK 7
|:|||||:
Db 236 QKLCQPK 242

RESULT 6
Q94164 PRELIMINARY; PRT; 208 AA.
AC Q94164;
DT 01-FEB-1997 (T-EMBLrel. 02, Created)
DT 01-FEB-1997 (T-EMBLrel. 02, Last sequence update)
DT 01-JUN-2001 (T-EMBLrel. 17, Last annotation update)
DE SIMILARITY TO BPTI/KUNITZ INHIBITOR DOMAIN.
CN C10G8.2.

OS Caenorhabditis elegans.
OC Eukaryota; Metazoa; Nematoda; Chromadorea; Rhabditida; Rhabditidae;
OC Rhabditidae; Pelodierinae; Caenorhabditis.
OX NCBI_TaxID=6239;
RN [1]
RP SEQUENCE FROM N.A.
RA STRAIN-BRISTOL N2;
RX MEDLINE=94150718; PubMed=7906398;
RA Wilson R., Ainscough R., Anderson K., Baynes C., Berks M.,
RA Bonfield J., Burton J., Connell M., Copsey T., Cooper J., Coulson A.,
RA Craxton M., Dear S., Du Z., Durbin R., Favell A., Fulton L.,
RA Gardner A., Green P., Hawkins T., Hillier L., Jier M., Johnston L.,
RA Jones M., Kershaw J., Kirsten J., Laister N., Latreille P.,
RA Lightning J., Lloyd C., McMurray A., Mortimore B., O'Callaghan M.,
RA Parsons J., Percy C., Rifkin L., Roopra A., Saunders D., Showkneen R.,
RA Snelton N., Smith A., Sonchamer E., Staden R., Sulston J.,
RA Thierry-Mieg J., Thomas K., Vaubin K., Vaughan K., Waterston R.,
RA Watson A., Weinstein L., Wilkinson-Sproat J., Wohlman P.;
RT "2.2 Mb of contiguous nucleotide sequence from chromosome III of C.
elegans.";
RL Nature 368:32-38(1994).
RN [2]
RP SEQUENCE FROM N.A.
RC STRAIN-BRISTOL N2;

RA Blanchard M., Bradshaw H.;
RL Submitted (OCT-1996) to the EMBL/GenBank/DBJ databases.
RN [3]
RP SEQUENCE FROM N.A.
RC STRAIN-BRISTOL N2;
RA Waterston R.;
RL Submitted (SEP-1996) to the EMBL/GenBank/DBJ databases.
CC -!- SIMILARITY: TO PANGREATIC TRYPsin INHIBITOR (KUNITZ) DOMAIN.
DR EMBL; U70857; AAB09170.1;
DR InterPro; IPR002223; Kunitz_BPTI.
DR Pfam; PF00014; Kunitz_BPTI; 1.
DR SMART; SM00131; KU; 1.
DR PROSITE; PS00280; BPTI_KUNITZ_1; 1.
DR PROSITE; PS0279; BPTI_KUNITZ_2; 1.
KW Serine protease inhibitor.
SQ SEQUENCE 208 AA; 24008 MW; 779ABA4A8948E67B0 CRC64;

Query Match 73.9%; Score 34; DB 5; Length 208;
Best Local Similarity 83.3%; Pred. No. 21;
Matches 5; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

QY 3 LCHQKK 8
|:|||||:
Db 172 MCHQKK 177

RESULT 7
Q9BSM2 PRELIMINARY; PRT; 322 AA.
AC Q9BSM2;
DT 01-JUN-2001 (T-EMBLrel. 17, Created)
DT 01-JUN-2001 (T-EMBLrel. 17, Last sequence update)
DT 01-JUN-2001 (T-EMBLrel. 17, Last annotation update)
DE UNKNOWN (PROTEIN FOR IMAGE:3619689) (FRAGMENT).
OS Homo sapiens (Human).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
OX NCBI_TaxID=9606;
RN [1]
RP SEQUENCE FROM N.A.
RA TISSUE=LYMPHOMA;
RA Strausberg R.;
RL Submitted (MAR-2001) to the EMBL/GenBank/DBJ databases.
DR EMBL; BC004950; AAH04950.1;
FT NON_TER 1
SQ SEQUENCE 322 AA; 34916 MW; 33C65E86S8F9D761 CRC64;

Query Match 73.9%; Score 34; DB 4; Length 322;
Best Local Similarity 71.4%; Pred. No. 29;
Matches 5; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

QY 1 QKLCQKK 7
|:|||||:
Db 274 QKICHSK 280

RESULT 8
Q9D3R6 PRELIMINARY; PRT; 409 AA.
AC Q9D3R6;
DT 01-JUN-2001 (T-EMBLrel. 17, Created)
DT 01-JUN-2001 (T-EMBLrel. 17, Last sequence update)
DT 01-JUN-2001 (T-EMBLrel. 17, Last annotation update)
DE 4933439B08RIK PROTEIN.
GN 4933439B08RIK.
OS Mus musculus (Mouse).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
OX NCBI_TaxID=10090;
RN [1]
RP SEQUENCE FROM N.A.

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RC STRAIN=C57BL/6J; TISSUE=TESTIS;
RX MEDLINE=21085660; PubMed=11217851;
RA Kawai J., Shinagawa A., Shibata K., Yoshino M., Itoh M., Ishii Y.,
RA Akawa T., Hara A., Fukunishi Y., Konno H., Aochi J., Fukuda S.,
RA Aizawa K., Izawa M., Nishi K., Kiyosawa H., Kondo S., Yamanaka I.,
RA Saito T., Okazaki Y., Gojohori T., Bono H., Kasukawa T., Saito R.,
RA Kadota K., Matsuda H.A., Ashburner M., Batalov S., Casavant T.,
RA Fleischmann W., Gaasterland T., Gissi C., King B., Kochiwa H.,
RA Kuehl P., Lewis S., Matsuo Y., Nikaido I., Pesole G., Quackenbush J.,
RA Schriml L.M., Stauble F., Suzuki R., Tomita M., Wagner L., Washio T.,
RA Sakai K., Okido T., Furuno M., Aono H., Baldarelli R., Barsh G.,
RA Blake J., Boffelli D., Bojunga N., Carninci P., de Bonaldo M.F.,
RA Brownstein M.J., Bult C., Fletcher C., Fujita M., Gariboldi M.,
RA Gustincich S., Hill D., Hofmann M., Hume D.A., Kaniya M., Lee N.H.,
RA Lyons P., Marchionni L., Mashima J., Mazzarelli J., Mombaerts P.,
RA Nordone P., Ring B., Schoenbach M., Rodriguez I., Sakamoto N.,
RA Sasaki H., Sato K., Schenbach C., Seva T., Shibata Y., Storch K.-F.,
RA Suzuki H., Toyooka K., Wang K.H., Weitz C., Whittaker C., Wilming L.,
RA Wyszynski-Boris A., Yoshida K., Hasegawa Y., Kawaji H., Kohsaki S.,
RA Hayashizaki Y.;
RT "Functional annotation of a full-length mouse cDNA collection.";
RL Nature 409:685-690(2001).
DR EMBL; AK017114; BAB30604.1; -.
DR MGD; MGI:1918456; 4933439B08R1K.
DR InterPro; IPR003593; AAA.
DR InterPro; IPR003593; AAA_subfam.
DR Pfam; PF00004; AAA; 1.
DR SMART; SM00382; AAA; 1.
DR SEQUENCE 409 AA; 46131 MW; 9600B2008DC9749 CRC64;

Query Match 73.9%; Score 34; DB 11; Length 409;
Best Local Similarity 83.3%; Pred No. 35;
Matches 5; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

QY 2 KICHOX 7
DB 122 KICHOX 127

RESULT 9
QH8F4 PRELIMINARY; PRT; 522 AA.
ID Q9H8F4
AC Q9H8F4;
DT 01-MAR-2001 (TrEMBLrel. 16, Created)
DT 01-MAR-2001 (TrEMBLrel. 16, Last sequence update)
DT 01-JUN-2001 (TrEMBLrel. 17, Last annotation update)
DE CDNA FLJ13679 FIS, CLONE PLAC2000006.
OS Homo sapiens (Human).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
OX NCBI_TaxID=9606;
RN [1]
RP SEQUENCE FROM N.A.
RC TISSUE=PLACENTA;
RA Isogai T., Ota T., Hayashi K., Sugiyama T., Otsuki T., Suzuki Y.,
RA Nishikawa T., Nagai K., Sugano S., Takahashi-Fujii A., Hara H.,
RA Tanase T., Nomura Y., Togiya S., Komai F., Hara R., Takeuchi K.,
RA Arita M., Nabekura T., Ishii S., Kawai Y., Saito K., Yamamoto J.,
RA Wakamatsu A., Nakamura Y., Nagahari K., Masuho Y., Oshima A.;
RT "NEO human cDNA sequencing project.";
RL Submitted (AUG-2000) to the EMBL/GenBank/DBJ databases.
DR EMBL; AK023741; BAB14563.1; -.
DR InterPro; IPR001849; PH.
DR InterPro; IPR000219; RhoGEF.
DR Pfam; PF00169; PH; 1.
DR Pfam; PF00521; RhoGEF; 1.
DR SMART; SM00233; PH; 1.
DR SMART; SM00325; RhoGEF; 1.
DR PROSITE; PSS0003; PH_DOMAIN; 1.
DR SEQUENCE 522 AA; 59222 MW; F01A6BF70D2920F9 CRC64;

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Query Match 73.9%; Score 34; DB 4; Length 522;
Best Local Similarity 62.5%; Pred. No. 42;
Matches 5; Conservative 3; Mismatches 0; Indels 0; Gaps 0;

QY 1 QKLCCHK 8
DB 157 QELCHQOE 164

RESULT 10
O74308 PRELIMINARY; PRT; 547 AA.
ID O74308
AC O74308;
DT 01-NOV-1998 (TrEMBLrel. 08, Created)
DT 01-NOV-1998 (TrEMBLrel. 08, Last sequence update)
DT 01-JUN-2001 (TrEMBLrel. 17, Last annotation update)
DE ZINC-FINGER PROTEIN.
GN SPC15D4.02.
OS Schizosaccharomyces pombe (Fission yeast).
OC Eukaryota; Fungi; Ascomycota; Schizosaccharomycetes;
OC Schizosaccharomycetales; Schizosaccharomycetaceae;
OC Schizosaccharomycetes.
OX NCBI_TaxID=4896;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=972H;
RA Lyne M., Rajadream M.A., Barrell B.G., Lucas M., Gaillardin C.;
RL Submitted (AUG-1997) to the EMBL/GenBank/DBJ databases.
CC -1- SUBCELLULAR LOCATION: NUCLEAR (BY SIMILARITY).
CC -1- SIMILARITY: CONTAINS A ZN(2)-CYS(6), FUNGAL-TYPE BINUCLEAR CLUSTER
CC DOMAIN.
DR EMBL; AL031349; CAA20477.1; -.
DR HSP; P25502; IZME.
DR InterPro; IPR001138; zn2_Cy6_fungal.
DR Pfam; PF00172; Zn_glu; 1.
DR SMART; SM00066; GAL4; 1.
DR PROSITE; PS00463; ZN2_Cy6_FUNGAL_1; 1.
DR PROSITE; PS00048; ZN2_Cy6_FUNGAL_2; 1.
KW DNA-binding; Metal-binding; Nuclear protein; Transcription regulation;
KW Zinc; Zinc-finger.
SQ SEQUENCE 547 AA; 59641 MW; A65PD7D039B6CD18 CRC64;

Query Match 73.9%; Score 34; DB 3; Length 547;
Best Local Similarity 71.4%; Pred. No. 43;
Matches 5; Conservative 2; Mismatches 0; Indels 0; Gaps 0;

QY 1 QKLCCHK 7
DB 427 KLCCHK 433

RESULT 11
Q9NVK9 PRELIMINARY; PRT; 790 AA.
ID Q9NVK9
AC Q9NVK9;
DT 01-OCT-2000 (TrEMBLrel. 15, Created)
DT 01-OCT-2000 (TrEMBLrel. 15, Last sequence update)
DT 01-JUN-2001 (TrEMBLrel. 17, Last annotation update)
DE CDNA FLJ10665 FTS, CLONE NT2Rp2006200.
OS Homo sapiens (Human)
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
OX NCBI_TaxID=9606;
RN [1]
RP SEQUENCE FROM N.A.
RA Isogai T., Ota T., Hayashi K., Sugiyama T., Otsuki T., Suzuki Y.,
RA Nishikawa T., Nagai K., Sugano S., Ishibashi T., Fujimori K.,
RA Tani H., Kimata M., Watanabe M., Hiraoka S., Ishii S., Kawai Y.,
RA Saito K., Yamamoto J., Wakamatsu A., Nakamura Y., Nagahari K.,
RA Masuho Y., Kanehori K.;
RT "NEO human cDNA sequencing project.";
RL Submitted (FEB-2000) to the EMBL/GenBank/DBJ databases.

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DR EMBL; AK001527; BAA91741.1; -
 DR InterPro; IPR001849; PH.
 DR InterPro; IPR000219; RhGEF.
 DR Pfam; PF00169; RhGEF.
 DR Pfam; PF00621; RhGEF; 1.
 DR SMART; SM00233; PH; 1.
 DR SMART; SM00325; RhGEF; 1.
 DR PROSITE; PS0003; PH_DOMAIN; 1.
 SQ SEQUENCE 790 AA; 88975 MW; A3B9F4972CE2D509 CRC64;

Query Match 73.9%; Score 34; DB 4; Length 790;
 Best Local Similarity 62.5%; Pred. No. 57;
 Matches 5; Conservative 3; Mismatches 0; Indels 0; Gaps 0;

QY 1 QKLCQK 8
 :||||:
 DB 157 QELCHQE 164

RESULT 12
 O96143 PRELIMINARY; PRT; 1188 AA.
 AC O96143;
 DT 01-MAY-1999 (TrEMBLrel. 10, Created)
 DT 01-MAY-1999 (TrEMBLrel. 10, Last sequence update)
 DT 01-MAY-1999 (TrEMBLrel. 10, Last annotation update)
 DE PROTEIN WITH 5'-3' EXONUCLEASE DOMAIN (KEM-1 FAMILY).
 GN PPB0205C.
 OS Plasmodium falciparum.
 OC Eukaryota; Alveolata; Apicomplexa; Haemosporidia; Plasmodium.
 OX NCBI_TaxID=5833;
 RN [1]
 RP SEQUENCE FROM N.A.
 RX MEDLINE-99021743; PubMed-9804551;
 RA Gardner M.J., Tetteelin H., Carucci D.J., Cummings L.M., Aravind L.,
 RA Koonin E.V., Shallom S., Mason T., Yu K., Fujii C., Pederson J.,
 RA Shen K., Jing J., Aston C., Lai Z., Schwartz D.C., Pertea M.,
 RA Salzberg S., Zhou L., Sutton G.G., Clayton R., White O., Smith H.O.,
 RA Fraser C.M., Adams M.D., Venter J.C., Hoffman S.L.;
 RT "Chromosome 2 sequence of the human malaria parasite Plasmodium
 falciparum".
 RL Science 282:1126-1132(1998).
 RW EMBL; AE001380; AAC71830.1; -
 KW Exonuclease.
 SQ SEQUENCE 1188 AA; 142895 MW; BF767FC8532EBAC9 CRC64;

Query Match 73.9%; Score 34; DB 5; Length 1188;
 Best Local Similarity 71.4%; Pred. No. 76;
 Matches 5; Conservative 2; Mismatches 0; Indels 0; Gaps 0;

QY 1 QKLCQK 7
 :||||:
 DB 1117 EELCHQK 1123

RESULT 13
 O13348 PRELIMINARY; PRT; 1295 AA.
 AC O13348;
 DT 01-JAN-1998 (TrEMBLrel. 05, Created)
 DT 01-JAN-1998 (TrEMBLrel. 05, Last sequence update)
 DT 01-JUN-2001 (TrEMBLrel. 17, Last annotation update)
 DE REVERSE TRANSCRIPTASE.
 OS Magnaporthe grisea (Rice blast fungus) (Pyricularia grisea).
 OC Eukaryota; Fungi; Ascomycota; Pezizomycotina; Sordariomycetes;
 OC Sordariomycetes incertae sedis; Magnaportheaceae; Magnaporthe.
 OX NCBI_TaxID=148305;
 RN [1]
 RP SEQUENCE FROM N.A.
 RX Meyn III M.A., Farrall L., Valent B., Chumley F.G., Orbach M.J.;
 RA Submitted (AUG-1997) to the EMBL/GenBank/DBJ databases.

CC -!- SIMILARITY: TO RNA-DIRECTED DNA POLYMERASE (REVERSE
 CC TRANSCRIPTASE).
 DR EMBL; AF018033; AAB71689.1; -
 DR InterPro; IPR002156; RNaseH.
 DR InterPro; IPR000477; RVTse.
 DR Pfam; PF00075; RNaseH; 1.
 DR Pfam; PF00078; RVT; 1.
 KW RNA-directed DNA polymerase.
 SQ SEQUENCE 1295 AA; 144307 MW; E811059B750D5421 CRC64;

Query Match 73.9%; Score 34; DB 3; Length 1295;
 Best Local Similarity 71.4%; Pred. No. 81;
 Matches 5; Conservative 2; Mismatches 0; Indels 0; Gaps 0;

QY 2 KLCHQK 8
 :||||:
 DB 33 ELCHQK 39

RESULT 14
 Q63625 PRELIMINARY; PRT; 1473 AA.
 AC Q63625;
 DT 01-NOV-1996 (TrEMBLrel. 01, Created)
 DT 01-NOV-1996 (TrEMBLrel. 01, Last sequence update)
 DT 01-NOV-1998 (TrEMBLrel. 08, Last annotation update)
 DE CTD-BINDING SR-LIKE PROTEIN RA9.
 OS Rattus norvegicus (Rat).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Rattus.
 OX NCBI_TaxID=10116;
 RN [1]
 RP SEQUENCE FROM N.A.
 RX TISSUE-HIPECAMPUS;
 RC MEDLINE-96293459; PubMed-8692929;
 RA Yuryev A., Patturajan M., Litingtung Y., Joshi R.V., Gentile C.,
 RA Gebara M., Corden J.L.;
 RT "The C-terminal domain of the largest subunit of RNA polymerase II
 interacts with a novel set of serine/arginine-rich proteins.";
 RL Proc. Natl. Acad. Sci. U.S.A. 93:6975-6980(1996).
 RW EMBL; U49057; AAC52858.1; -
 SQ SEQUENCE 1473 AA; 161203 MW; 949EE6F5873989BF CRC64;

Query Match 73.9%; Score 34; DB 11; Length 1473;
 Best Local Similarity 71.4%; Pred. No. 89;
 Matches 5; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

QY 1 QKLCQK 7
 :||||:
 DB 1425 QKCHSK 1431

RESULT 15
 Q9PLY6 PRELIMINARY; PRT; 1654 AA.
 AC Q9PLY6;
 DT 01-OCT-2000 (TrEMBLrel. 15, Created)
 DT 01-OCT-2000 (TrEMBLrel. 15, Last sequence update)
 DT 01-JUN-2001 (TrEMBLrel. 17, Last annotation update)
 DE KIAA1542 PROTEIN (FRAGMENT).
 GN KIAA1542
 OS Homo sapiens (Human).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 OC Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
 OX NCBI_TaxID=9606;
 RN [1]
 RP SEQUENCE FROM N.A.
 RX MEDLINE-20277482; PubMed-10819331;
 RA Nagase T., Kikuno R., Ishikawa K., Hirose M., Ohara O.;
 RT "Prediction of the coding sequences of unidentified human
 genes. XVII. The complete sequences of 100 new cDNA clones from brain

RT which code for large proteins in vitro.
RL DNA Res. 7:143-150(2000).
CC -!- SIMILARITY: CONTAINS A RING-TYPE ZINC FINGER.
DR EMBL; AB040975; BAA96066.1; -
DR InterPro; IPR002219; DAG_PE-bind.
DR InterPro; IPR001965; PHD_PE-bind.
DR InterPro; IPR001841; Znf_ring.
DR Pfam; PF00628; PHD; 1.
DR SMART; SM00109; CL; 1.
DR SMART; SM00249; PHD; 1.
DR SMART; SM00184; RING; 2.
DR PROSITE; PS00518; ZINC_FINGER_C3HC4; 1.
KW Zinc-finger.
FT NON_TER 1
SQ SEQUENCE 1654 AA; 179053 MW; 125CF71A84AFB218 CRC64;

Query Match 73.9%; Score 34; DB 4; Length 1654;
Best Local Similarity 71.4%; Pred. No. 97;
Matches 5; Conservative 1; Mismatches 1; Indels 0; Gaps 0;
QY 1 QKLCHQK 7
DB 1606 QXCHSK 1612

Search completed: February 12, 2002, 12:03:26
Job time: 817 sec

GenCore version 4.5
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OM protein - protein search, using sw model

Run On: February 12, 2002, 12:04:02 ; Search time 30.28 seconds
(without alignments)
9.887 Million cell updates/sec

Title: US-09-606-129A-19
Perfect score: 46
Sequence: 1 QKLCHQKK 8

Scoring table: BLOSUM62
Gapop 10.0 , Gapext 0.5

Searched: 100059 seqs, 36664827 residues
Total number of hits satisfying chosen parameters: 100059

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database : SwissProt_39.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match %	Length	DB ID	Description
1	46	100.0	295	1 BIEA_RAT	P46844 rattus norv
2	36	78.3	654	1 CBP1_YEAST	P07252 saccharomyc
3	33	71.7	673	1 Z145_HUMAN	Q05516 homo sapien
4	32	69.6	135	1 VE6_PAPVE	P11331 european el
5	32	69.6	210	1 YQ33_CAEEL	Q09311 caenorhabdi
6	32	69.6	272	1 Q3HQ_ACICA	Q59087 acinetobact
7	32	69.6	345	1 TF2B_YEAST	P29055 saccharomyc
8	32	69.6	357	1 Y303_MYCGE	P47545 mycoplasma
9	32	69.6	595	1 PRIM_CHLTR	O84799 chlamydia t
10	32	69.6	609	1 FETA_GORGO	P28050 gorilla gor
11	32	69.6	609	1 FETA_HUMAN	P02771 homo sapien
12	32	69.6	645	1 YQ81_CAEEL	P34617 caenorhabdi
13	32	69.6	595	1 SY1_MYCGE	P47587 mycoplasma
14	31	67.4	75	1 EX7S_BACSU	P54522 bacillus su
15	31	67.4	169	1 NEUT_BOVIN	P01156 bos taurus
16	31	67.4	170	1 NEUT_CANFA	P10673 canis famil
17	31	67.4	271	1 YSM4_CAEEL	Q10124 caenorhabdi
18	31	67.4	478	1 ARDE_CHLTR	O84375 chlamydia t
19	31	67.4	498	1 NFSL_SCHPO	O74351 schizosacch
20	31	67.4	677	1 NRGL_XENLA	O93383 xenopus lae
21	31	67.4	892	1 LD12_XENLA	Q99088 xenopus lae
22	31	67.4	1207	1 DPOL_ASFB7	P42489 african swi
23	31	67.4	1244	1 DPOL_ASFL6	P43139 african swi
24	31	67.4	1256	1 FLII_DROME	Q24020 drosophila
25	31	67.4	1376	1 MYHA_BOVIN	Q27991 bos taurus
26	31	67.4	1376	1 MYHA_HUMAN	P35580 homo sapien
27	31	67.4	1376	1 MYHA_RAT	Q9J100 rattus norv
28	30	65.2	80	1 EX7S_BACHD	Q9K968 bacillus ha
29	30	65.2	159	1 MPAA_CORAV	Q08407 coryllus ave
30	30	65.2	259	1 DEOC_ECOLI	P00882 escherichia
31	30	65.2	347	1 YF86_MYCPN	P75194 mycoplasma
32	30	65.2	397	1 CAPB_BACAN	P19580 bacillus an
33	30	65.2	442	1 GAG_VILV	P03352 visna lenti

34	30	65.2	442	1 GAG_VILV1	P23424 visna lenti
35	30	65.2	442	1 GAG_VILV2	P23425 visna lenti
36	30	65.2	442	1 GAG_VILV3	P35955 visna lenti
37	30	65.2	446	1 GAG-OMVVS	P16900 ovine lenti
38	30	65.2	457	1 LAT_STRCL	Q01767 streptomyce
39	30	65.2	489	1 OCLN_POTTR	Q28793 potorous tr
40	30	65.2	583	1 HASS_XENLA	O57428 xenopus lae
41	30	65.2	611	1 ACE_HAEIE	Q10715 haematobia
42	30	65.2	686	1 TRFE-ANAPL	P56410 anas platyr
43	30	65.2	705	1 TRFE-CHICK	P02789 gallus gall
44	30	65.2	753	1 YP6A_CAEEL	Q09219 caenorhabdi
45	30	65.2	837	1 SM4G_MOUSE	Q9whu7 mus musculu

ALIGNMENTS

RESULT 1
BIEA_RAT ID BIEA_RAT STANDARD: PRT: 295 AA.
AC P46844;
DT 01-NOV-1995 (Rel. 32, Created)
DT 01-NOV-1995 (Rel. 32, Last sequence update)
DT 15-JUL-1993 (Rel. 38, Last annotation update)
DE BILIVERDIN REDUCTASE A PRECURSOR (EC 1.3.1.24) (BILIVERDIN-IX ALPHA-REDUCTASE).
GN BLVRA OR BLVR.
OS Rattus norvegicus (Rat).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Rattus.
OX NCBI_TaxID=10116;
RN [1]
RP SEQUENCE FROM N.A., AND PARTIAL SEQUENCE.
RC TISSUE=Kidney;
RX MEDLINE=92156147; PubMed=1371282;
RA Fakhrai H., Maines M.D.;
RT "Expression and characterization of a cDNA for rat kidney biliverdin reductase. Evidence suggesting the liver and kidney enzymes are the same transcript product.";
RL J. Biol. Chem. 267:4023-4029(1992).
RN [2]
RX MUTAGENESIS.
RX MEDLINE=94291657; PubMed=8020496;
RA McCoubrey W.K. Jr., Maines M.D.;
RT "Site-directed mutagenesis of cysteine residues in biliverdin reductase. Roles in substrate and cofactor binding.";
RL Eur. J. Biochem. 222:597-603(1994).
CC -!- FUNCTION: CONVERTS BILIVERDIN TO BILIRUBIN; DISPLAYS TWO DISTINCT PH OPTIMA USING A DIFFERENT COFACTOR AT EACH PH: NADH AT THE LOWER PH 6.7-6.9 RANGE AND NADPH AT PH 8.5-8.7. NADPH, HOWEVER, IS THE PROBABLE COFACTOR IN BIOLOGICAL SYSTEMS.
CC -!- CATALYTIC ACTIVITY: BILIRUBIN + NAD(P)(+) -> BILIVERDIN + NAD(P)H.
CC -!- COFACTOR: BINDS ONE ZINC ION.
CC -!- PATHWAY: FINAL STEP IN HEME METABOLISM.
CC -!- SUBUNIT: MONOMER (BY SIMILARITY).
CC -!- SUBCELLULAR LOCATION: CYTOPLASMIC.
CC -!- SIMILARITY: TO E.COLI YHHX.
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CC -----
CC EMBL; M81681; AAA40830.1; -
CC Interpro; IPR000683; GFO_IDH_MoCA.
CC Pfam; PF01408; GFO_IDH_MoCA; 1.
CC Oxidoreductase; NAD; NADP; Zinc.
CC PROPEP 1 2
CC CHAIN 3 295 BILIVERDIN REDUCTASE A.
CC DOMAIN 11 16 POLY-VAL.

FT METAL 279 ZINC (POTENTIAL).
FT METAL 280 ZINC (POTENTIAL).
FT METAL 291 ZINC (POTENTIAL).
FT METAL 292 ZINC (POTENTIAL).
FT MUTAGEN 73 C->A: LOSS OF ACTIVITY.
FT MUTAGEN 280 C->A: REDUCED ACTIVITY.
FT MUTAGEN 291 C->A: REDUCED ACTIVITY.
SQ SEQUENCE 295 AA; 33565 MW; 219C8EA95C150588 CRC64;

Query Match 100.0%; Score 46; DB 1; Length 295;
Best Local Similarity 100.0%; Pred. No. 0.18; Indels 0; Gaps 0;
Matches 8; Conservative 0; Mismatches 0;

QY 1 QKLCHQKK 8
DB 288 QKLCHQKK 295

RESULT 2
CBPL_YEAST STANDARD; PRT; 654 AA.
AC P07252;
DT 01-APR-1988 (Rel. 07, Created)
DT 01-APR-1988 (Rel. 07, Last sequence update)
DT 01-NOV-1987 (Rel. 35, Last annotation update)
DE CYTOCHROME B PRE-MRNA PROCESSING PROTEIN 1.
GN CBPI OR YJL205W OR J0242 OR HRA634.
OS Saccharomyces cerevisiae (Baker's yeast).
OC Eukaryota; Fungi; Ascomycota; Saccharomycotina; Saccharomycetes;
OC Saccharomycetales; Saccharomycetaceae; Saccharomycetes.
OX NCBI_TaxID=4932;
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE=84185566; PubMed=6325407;
RA Dieckmann C.L., Homison G., Tzagoloff A.;
RT "Assembly of the mitochondrial membrane system. Nucleotide sequence of a yeast nuclear gene (CBP1) involved in 5' end processing of cytochrome b pre-mRNA";
RL J. Biol. Chem. 259:4732-4738(1984).
RN [2]
RP SEQUENCE FROM N.A.
RX STRAIN=S288C;
RA Vandenberg M., Durand P., Bolle P.-A., Dion C., Portetelle D., Hilger F.;
RT "Sequence analysis of a 40.2 kb DNA fragment located near the left telomere of yeast chromosome X";
RL Yeast 10:1657-1662(1994).
RN [3]
RP SEQUENCE OF 589-654 FROM N.A.
RX MEDLINE=90014786; PubMed=2552292;
RA Liu Y., Dieckmann C.L.;
RT "Overproduction of yeast viruslike particles by strains deficient in a mitochondrial nuclease";
RL Mol. Cell. Biol. 9:3323-3331(1989).
CC -!- FUNCTION: RESPONSIBLE FOR CONFERRING A STABLE 5' END ON CYTOCHROME B MRNA.

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CC EMBL: K02647; AAA34474.1;
CC EMBL: Z34098; CA884002.1;
CC EMBL: Z49484; CA889506.1;
CC EMBL: M28067; AAA34456.1;
CC PIR: S05829; BVBP1.
CC PIR: S45164; S45164.

DR SGD; S0003745; CBP1.
KW mRNA processing.
SQ SEQUENCE 654 AA; 76171 MW; 2453B03280E1044D CRC64;

Query Match 78.3%; Score 36; DB 1; Length 654;
Best Local Similarity 75.0%; Pred. No. 21;
Matches 6; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

QY 1 QKLCHQKK 8
DB 514 KKLCHYKK 521

RESULT 3
Z145_HUMAN STANDARD; PRT; 673 AA.
ID Z145_HUMAN
AC Q05516;
DT 01-NOV-1995 (Rel. 32, Created)
DT 01-NOV-1995 (Rel. 32, Last sequence update)
DT 20-AUG-2001 (Rel. 40, Last annotation update)
DE ZINC FINGER PROTEIN PLZF (PROMYELOCYTIC LEUKEMIA ZINC FINGER DE PROTEIN) (ZINC FINGER PROTEIN 145).
GN ZNF145 OR PLZF.
OS Homo sapiens (Human).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
OX NCBI_TaxID=9606;
RN [1]
RP SEQUENCE FROM N.A.
RC TISSUE=Heart ventricle;
RX MEDLINE=93209216; PubMed=8384553;
RA Chen Z., Brand N.J., Chen A., Chen S.-J., Tong J.-H., Wang Z.-Y., Waxman S., Zelen A.;
RT "Fusion between a novel Kruppel-like zinc finger gene and the retinoic acid receptor-alpha locus due to a variant t(11;17) translocation associated with acute promyelocytic leukaemia";
RL EMBO J. 12:1161-1167(1993).
RN [2]
RP SEQUENCE OF 424-455 FROM N.A.
RX MEDLINE=93253074; PubMed=8387545;
RA Chen S.-J., Zelen A., Tong J.-H., Yu H.-Q., Wang Z.-Y., Derre J., Berger R., Waxman S., Chen Z.;
RT "Rearrangements of the retinoic acid receptor alpha and promyelocytic leukemia zinc finger genes resulting from t(11;17)(q23;q21) in a patient with acute promyelocytic leukemia";
RL J. Clin. Invest. 91:2260-2267(1993).
RN [3]
RP X-RAY CRYSTALLOGRAPHY (2.0 ANGSTROMS) OF 7-122.
RX MEDLINE=20005701; PubMed=10537309;
RA Li X., Peng H., Schultz D.C., Lopez-Guisa J.M., Rauscher F.J. III, Marmorstein R.;
RT "Structure-function studies of the BTB/POZ transcriptional repression domain from the promyelocytic leukemia zinc finger oncoprotein";
RL Cancer Res. 59:5275-5282(1999).
CC -!- FUNCTION: PROBABLE TRANSCRIPTION FACTOR. MAY PLAY A ROLE IN MYELOID MATURATION AND IN THE DEVELOPMENT AND/OR MAINTENANCE OF OTHER DIFFERENTIATED TISSUES.
CC -!- SUBCELLULAR LOCATION: NUCLEAR.
CC -!- ALTERNATIVE PRODUCTS: 2 ISOFORMS; PLZF AND PLZF2 (SHOWN HERE); ARE PRODUCED BY ALTERNATIVE SPLICING.
CC -!- TISSUE SPECIFICITY: WITHIN THE HEMATOPOIETIC SYSTEM, PLZF IS EXPRESSED IN BONE MARROW, EARLY MYELOID CELL LINES AND PERIPHERAL BLOOD MONONUCLEAR CELLS. ALSO EXPRESSED IN THE OVARY, AND AT LOWER LEVELS, IN THE KIDNEY AND LUNG.
CC -!- INDUCTION: BY RETINOIC ACID.
CC -!- DISEASE: A FORM OF ACUTE PROMYELOCYTIC LEUKEMIA (APL) IS CHARACTERIZED BY A CHROMOSOMAL TRANSLOCATION T(11;17)(Q32;Q21) WHICH INVOLVES ZNF145 AND RETINOIC ACID RECEPTOR ALPHA (RARA).
CC -!- SIMILARITY: BELONGS TO THE KRUEPPEL FAMILY OF C2H2-TYPE ZINC-FINGER PROTEINS.
CC -!- SIMILARITY: CONTAINS 1 BTB/POZ DOMAIN.

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CC or send an email to license@isb-sib.ch).
CC -----
DR EMBL; Z19002; CAA79489.1; -
DR EMBL; S60093; AAC60590.2; -
DR PDB; 1CS3; 09-AUG-00.
DR TRANSFAC; T02336; -
DR MIM; 176797; -
DR InterPro; IPR000210; BTB_POZ.
DR InterPro; IPR000822; Znf-C2H2.
DR Pfam; PF00651; BTB; 1.
DR Pfam; PF00096; zf-C2H2; 9.
DR PRINTS; PR00048; ZINC_FINGER.
DR SMART; SM00225; BTB; 1.
DR SMART; SM00355; Znf_C2H2; 9.
DR PROSITE; PS00097; BTB; 1.
DR PROSITE; PS00028; ZINC_FINGER_C2H2_1; 8.
DR PROSITE; PS0157; ZINC_FINGER_C2H2_2; 9.
KW Transcription regulation; DNA-binding; Zinc-finger; Metal-binding;
KW Nuclear protein; Repeat; Chromosomal translocation; Proto-oncogene;
KW Phosphorylation; Alternative splicing; 3D-structure.
FT DOMAIN 34 96
FT MOD_RES 404 652
FT ZN_FING 404 426
FT ZN_FING 432 454
FT ZN_FING 461 483
FT ZN_FING 490 512
FT ZN_FING 518 540
FT ZN_FING 546 568
FT ZN_FING 574 596
FT ZN_FING 602 624
FT ZN_FING 630 652
FT SITE 394 395
FT BREAKPOINT FOR TRANSLOCATION TO FORM
FT PLZF-RAR-ALPHA ONCOGENE.
FT PHOSPHORYLATION (BY PDPK) (POTENTIAL).
FT MOD_RES 184 184
FT MOD_RES 197 197
FT MOD_RES 256 256
FT MOD_RES 282 282
FT MOD_RES 628 628
FT MOD_RES 655 377
FT VARSPLIC 255 377
FT MISSING (IN ISOFORM PL2FA).
FT SEQUENCE 673 AA; 74332 MW; 7CD7319E2A32109D CRC64;

Query Match 71.7%; Score 33; DB 1; Length 673;
Best Local Similarity 83.3%; Pred. No. 75;
Matches 5; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

QY 2 KLCQK 7
DB 605 KLCQK 510
|||||

RESULT 4
VE6_PAPVE STANDARD; PRT; 135 AA.
AC P11331;
DT 01-JUL-1989 (Rel. 11, Created)
DT 01-JUL-1989 (Rel. 11, Last sequence update)
DT 01-JUN-1994 (Rel. 29, Last annotation update)
DE E6 PROTEIN.
GN E6.
OS European elk papillomavirus (E6P).
OC Viruses; dsDNA viruses, no RNA stage; Papillomaviridae;
OC Papillomavirus.
OX NCBI_TaxID=10565;
RN [1]
RP SEQUENCE FROM N.A.
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RX MEDLINE=87219878; PubMed=3034730;
RA Ahola H., Bergman P., Stroem A.C., Moreno-Lopez J., Petterson U.;
RT "Organization and expression of the transforming region from the
RT European elk papillomavirus (E6P).";
RL Gene 50:195-205(1986).
CC -!- FUNCTION: EXHIBIT A STRONG, BUT NON SPECIFIC AFFINITY FOR DOUBLE
CC -!- SUBCELLULAR LOCATION: NUCLEAR MATRIX-ASSOCIATED.
CC -----
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CC -----
DR EMBL; M15953; AAA66849.1; -
DR PIR; A29499; W6WLEP. E6.
DR InterPro; IPR001334; E6.
DR Pfam; PF00518; E6; 1.
DR Early protein; DNA-binding; Nuclear protein; Zinc-finger.
FT ZN_FING 11 47
FT ZN_FING 83 119
FT POTENTIAL.
FT SEQUENCE 135 AA; 15869 MW; AE4F1BABC95E0459 CRC64;

Query Match 69.6%; Score 32; DB 1; Length 135;
Best Local Similarity 71.4%; Pred. No. 28;
Matches 5; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

QY 2 KLCQK 8
DB 24 KRCHEK 30
|||||

RESULT 5
YQS3_CAEEL STANDARD; PRT; 210 AA.
AC Q09311;
DT 01-NOV-1997 (Rel. 35, Created)
DT 01-NOV-1997 (Rel. 35, Last sequence update)
DT 01-NOV-1997 (Rel. 35, Last annotation update)
DE HYPOTHETICAL 23.6 KDA PROTEIN F21H12.3 IN CHROMOSOME II.
GN F21H12.3.
OS Caenorhabditis elegans.
OC Rhabditida; Metazoa; Nematoda; Chromadorea; Rhabditida; Rhabditoides;
OC Rhabditidae; Telodermata; Caenorhabditis.
OX NCBI_TaxID=6239;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=BRISTOL N2;
RA Pavello T.;
RL Submitted (JUL-1995) to the EMBL/GenBank/DBJ databases.
CC -!- SIMILARITY: TO C.ELEGANS ZK675.1.
CC -----
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CC -----
DR EMBL; U23176; AAC46715.1; -
DR WormPep; F21H12.3; CE01914.
KW Hypothetical protein.
FT SEQUENCE 210 AA; 23617 MW; 1E646E6FC30154A0 CRC64;

Query Match 69.6%; Score 32; DB 1; Length 210;
Best Local Similarity 71.4%; Pred. No. 41;
Matches 5; Conservative 1; Mismatches 1; Indels 0; Gaps 0;
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QY 1 QKLCQK 7
Db 23 KNLCHQK 29

RESULT 6
3DHQ_ACICA STANDARD; PRT; 272 AA.
AC Q59087;
DT 20-AUG-2001 (Rel. 40, Created)
DT 20-AUG-2001 (Rel. 40, Last sequence update)
DT 20-AUG-2001 (Rel. 40, Last annotation update)
DE CATABOLIC 3-DEHYDROQUINATE DEHYDRATASE (EC 4.2.1.10) (3-
DE DEHYDROQUINASE).
GN QUIB.
OS Acinetobacter calcoaceticus.
OC Bacteria; Proteobacteria; gamma subdivision; Moraxellaceae;
OC Acinetobacter.
OX NCBI_TaxID=471;
RN [1]
RP SEQUENCE FROM N.A., AND CHARACTERIZATION.
RC STRAIN=BD413 / ADP1;
RX MEDLINE=96011389; PubMed=7592351;
RA Elsenmore D.A., Ornstun L.N.;
RT "Unusual ancestry of dehydratases associated with quinate catabolism
RT in Acinetobacter calcoaceticus";
RL J. Bacteriol. 177:5971-5978(1995).
CC -!- FUNCTION: CATALYZES THE CATABOLIC DEHYDRATATION OF DEHYDROQUINATE
CC TO DEHYDROSHIKIMATE.
CC -!- CATALYTIC ACTIVITY: 3-DEHYDROQUINATE + H(2)O.
CC -!- PATHWAY: QUINIC ACID CATABOLIC PATHWAY; SECOND STEP. THIS PATHWAY
CC ALLOWS GROWTH OF BACTERIA WITH QUINATE BY ITS CONVERSION TO
CC PROTOCATECHUATE AND SUBSEQUENT METABOLISM BY THE BETA-KETOADIPATE
CC PATHWAY.
CC -!- INDUCTION: BY PROTOCATECHUATE.
CC -!- SIMILARITY: BELONGS TO THE TYPE-I 3-DEHYDROQUINASE FAMILY.
CC
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CC
CC EMBL: L05770; AAC37158.1; ALT_INIT.
CC InterPro: IPR001381; DHQinase_I.
CC Pfam: PF01487; DHQinase_I; 1.
CC PROSITE: PS01028; DEHYDROQUINASE_I; 1.
CC Quinate metabolism; Lyase.
KW ACT_SITE 163 190 BY SIMILARITY.
FT ACT_SITE 190 190 FORMS A SCHIFF-BASE INTERMEDIATE
FT REPEAT 272 345 (BY SIMILARITY).
FT SEQUENCE 272 AA; 29899 MW; 98646DC5E88BF6D3 CRC64;

Query Match 69.6%; Score 32; DB 1; Length 272;
Best Local Similarity 85.7%; Pred. NO. 51;
Matches 6; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 2 KLCQK 8
Db 149 KLAHQK 155

RESULT 7
TF2B_YEAST STANDARD; PRT; 345 AA.
AC P29055;
DT 01-DEC-1992 (Rel. 24, Created)
DT 01-DEC-1992 (Rel. 24, Last sequence update)
DT 15-JUL-1998 (Rel. 36, Last annotation update)

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DE DE
GN SUA7 OR YPR086W OR P9513.4.
OS Saccharomyces cerevisiae (Baker's yeast).
OC Eukaryota; Fungi; Ascomycota; Saccharomycotina; Saccharomycetes;
OC Saccharomycetales; Saccharomycetaceae; Saccharomycetes.
OX NCBI_TaxID=4932;
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE=92191276; PubMed=1547497;
RA Pinto I., Ware D.E., Hampsey M.;
RT "The yeast SUA7 gene encodes a homolog of human transcription factor
RT TFIIIB and is required for normal start site selection in vivo.";
RL Cell 68:977-988(1992).
RN [2]
RP SEQUENCE FROM N.A.
RC STRAIN=S288C / AB972;
RA Johnston M., Andrews S., Brinkman R., Cooper J., Ding H., Du Z.,
RA Favella A., Fulton L., Gattung S., Greco T., Kirsten J.,
RA Kucaba T., Hallsworth K., Hawkins J., Hillier L., Jier M.,
RA Johnson D., Johnston L., Langston Y., Latreille P., Le T.,
RA Mardis E., Meneses S., Miller N., Nian M., Pauley A., Peluso D.,
RA Rifken L., Riles L., Taich A., Trevaskis E., Vignati D.,
RA Wilcox L., Wohlman P., Vaudin M., Wilson R., Waterston R.;
RL Submitted (MAR-1996) to the EMBL/GenBank/DBJ databases.
CC -!- FUNCTION: GENERAL FACTOR THAT PLAYS A MAJOR ROLE IN THE ACTIVATION
CC OF EUKARYOTIC GENES TRANSCRIBED BY RNA POLYMERASE II.
CC -!- SUBUNIT: ASSOCIATES WITH TFIIID-11A (DA COMPLEX) TO FORM TFIIID-
CC 11A-11B (DAB-COMPLEX) WHICH IS THEN RECOGNIZED BY POLYMERASE II.
CC -!- SUBCELLULAR LOCATION: NUCLEAR.
CC -!- SIMILARITY: BELONGS TO THE TFIIIB FAMILY.
CC
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CC
CC EMBL: M81380; AAA35126.1; -.
CC EMBL: U51033; AAB68135.1; -.
CC PIR: S26707; S26707.
CC HSP: Q00403; 1TFB.
CC TRANSFAC: T00819; -.
CC SGD: S0006290; SUA7.
CC InterPro: IPR000553; Cyclin.
CC Pfam: PF00382; transcript_fac2; 2.
CC PRINTS: PS00685; TIFACTORIIB.
CC SMART: SM00385; CYCLIN; 2.
CC PROSITE: PS00782; TFIIIB; 1.
KW Transcription regulation; Nuclear protein; Repeat; Zinc-finger.
FT ZN_FING 24 48 POTENTIAL.
FT REPEAT 136 212
FT REPEAT 242 318
FT SEQUENCE 345 AA; 38200 MW; 8F1F6D24602436E2 CRC64;

Query Match 69.6%; Score 32; DB 1; Length 345;
Best Local Similarity 71.4%; Pred. NO. 63;
Matches 5; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

QY 2 KLCQK 8
Db 155 KLCHEK 161

RESULT 8
Y303_MYCGE STANDARD; PRT; 357 AA.
ID Y303_MYCGE
AC P47545;
DT 01-OCT-1996 (Rel. 34, Created)
DT 01-OCT-1996 (Rel. 34, Last sequence update)

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DT 20-AUG-2001 (Rel. 40, Last annotation update)
DE HYPOTHETICAL ABC TRANSPORTER ATP-BINDING PROTEIN MG303.
GN MG303.
OS Mycoplasma genitalium.
OC Bacteria; Firmicutes; Bacillus/Clostridium group; Mollicutes;
CC Mycoplasmataceae; Mycoplasma.
OX NCBI_TaxID=2097;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=ATCC 33530 / G-37;
RA Fraser C.M., Gocayne J.D., White O., Adams M.D., Clayton R.A., Kelley J.M.,
RA Fleischmann R.D., Bult C.J., Kerlavage A.R., Sutton G., Fuhrmann J.L.,
RA Fritchman J.L., Weidman J.F., Small K.V., Sandusky M., Phillips C.A., Merrick J.M.,
RA Nguyen D.T., Utterback T.R., Saudek D.M., Hu P.-C., Lucier T.S.,
RA Tomb J.F., Dougherty B.A., Bott K.F., Hutchison C.A. III, Venter J.C.,
RA Peterson S.N., Smith H.O., Hutchison C.A. III, Venter J.C.;
RT "The minimal gene complement of Mycoplasma genitalium.";
RL Science 270:397-403(1995).
CC -1- SIMILARITY: BELONGS TO THE ATP-BINDING TRANSPORT PROTEIN FAMILY
CC (ABC TRANSPORTERS).
CC -----
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CC -----
CC EMBL; U39710; AAC71525.1; -.
CC TIGR; MG303; -.
CC InterPro; IPR003593; AAA.
CC InterPro; IPR003439; ABC_transportr.
CC InterPro; IPR001687; ATP_GTP_A.
CC Pfam; PF00005; ABC_tran; 1.
CC SMART; SM00382; AAA; 1.
CC PROSITE; PS00211; ABC_TRANSPORTER; FALSE_NEG.
CC Hypothetical protein; ATP-binding; Transport; Complete proteome.
KW NP_BIND 107 114 ATP (POTENTIAL)
FT SEQUENCE 357 AA; 40786 MW; AFB1012F868E090E CRC64;
SQ
Query Match 69.6%; Score 32; DB 1; Length 357;
Best Local Similarity 83.3%; Pred. No. 65;
Matches 5; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

Qy' 2 KLCHK 7
Db 183 KLCHK 188
|||||
PRIM_CHLTR STANDARD; PRT; 595 AA.
AC 084799;
DT 20-AUG-2001 (Rel. 40, Created)
DT 20-AUG-2001 (Rel. 40, Last sequence update)
DT 20-AUG-2001 (Rel. 40, Last annotation update)
DE DNA PRIMASE (EC 2.7.7.-).
DN DNG OR CT794.
OS Chlamydia trachomatis.
OC Bacteria; Chlamydiales; Chlamydiaceae; Chlamydia.
OX NCBI_TaxID=813;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=D/UV-3/CX;
RA Stephens R.S., Kalman S., Lammel C.J., Fan J., Marathe R., Aravind L.,
RA Mitchell W.P., Olinger L., Tatusov R.L., Zhao Q., Koonin E.V.,
RA Davis R.W.;
RT "Genome sequence of an obligate intracellular pathogen of humans:
RT Chlamydia trachomatis.";

Science 282:754-759(1998).
-1- FUNCTION: DNA PRIMASE IS THE POLYMERASE THAT SYNTHESIZES SMALL
RNA PRIMERS FOR THE OKAZAKI FRAGMENTS ON BOTH TEMPLATE STRANDS AT
REPLICATION FORKS DURING CHROMOSOMAL DNA SYNTHESIS.
-1- COFACTOR: BINDS ONE ZINC ION PER MOLECULE (BY SIMILARITY).
-1- SUBUNIT: MONOMER (BY SIMILARITY).
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-----
EMBL; AE001351; AAC68389.1; -.
InterPro; IPR002936; Toprim.
InterPro; IPR002694; Znf_CHCC.
Pfam; PF01751; Toprim; 1.
Pfam; PF01807; Zf-CHC2; 1.
ProDom; PD002988; Znf_CHCC; 1.
SMART; SM00493; TOPRIM; 1.
SMART; SM00400; Znf_CHCC; 1.
KW Transferase; DNA replication; DNA-directed RNA polymerase; Primosome;
Zinc; Metal-binding; Complete proteome.
FT ZN_FING 39 622 CHC2-TYPE (BY SIMILARITY).
SQ SEQUENCE 595 AA; 58037 MW; 536858EBAFCD8FB6 CRC64;

Query Match 69.6%; Score 32; DB 1; Length 595;
Best Local Similarity 83.3%; Pred. No. 1e+02;
Matches 5; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

Qy 1 QKLCHQ 6
Db 565 RKLCHQ 570
|||||
FETA_GORGO STANDARD; PRT; 609 AA.
AC P28050;
DT 01-AUG-1992 (Rel. 23, Created)
DT 01-AUG-1992 (Rel. 23, Last sequence update)
DT 20-AUG-2001 (Rel. 40, Last annotation update)
DE ALPHA-FETOPROTEIN PRECURSOR (ALPHA-FETOGLOBULIN) (ALPHA-1-
DE FETOPROTEIN).
GN APP.
OS Gorilla gorilla gorilla (Lowland gorilla).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Gorilla.
OX NCBI_TaxID=9595;
RN [1]
RP SEQUENCE FROM N.A.
RC MEDLINE=91169517; PubMed=1706310;
RA Ryan S.C., Zielinski R., Dugaiczak A.;
RT "Structure of the gorilla alpha-fetoprotein gene and the divergence
RT of primates.";
RL Genomics 9:60-72(1991).
CC -1- FUNCTION: BINDS COPPER, NICKEL, AND FATTY ACIDS AS WELL AS, AND
CC BILIRUBIN LESS WELL THAN, SERUM ALBUMIN.
CC -1- SUBUNIT: DIMERIC AND TRIMERIC FORMS HAVE BEEN FOUND IN ADDITION
CC TO THE MONOMERIC FORM (BY SIMILARITY).
CC -1- SUBCELLULAR LOCATION: EXTRACELLULAR.
CC -1- TISSUE SPECIFICITY: PLASMA.
CC -1- DOMAIN: COMPOSED OF THREE HOMOLOGOUS DOMAINS.
CC -1- PTM: SULFATED (BY SIMILARITY).
CC -1- SIMILARITY: BELONGS TO THE ALB/APP/VDB FAMILY.
-----
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EMBL; M38272; AAA73520.1; "

PIR; A37970; FPGO.

HSSP; P02768; IUOR.

InterPro; IPR000264; Serum_albumin.

PIR; PF00273; transport_prot; 3.

PRINTS; PR00802; SERUMALBUMIN.

PRINTS; PR00803; AFETOPROTEIN.

SMART; SM00103; ALBUMIN; 3.

PROSITE; PS00212; ALBUMIN; 2.

Glycoprotein; Sulfation; Albumin; Plasma; Embryo; Repeat;

Metal-binding; Copper; Nickel; Signal.

FT SIGNAL 1 18

BY SIMILARITY.

ALPHA-FETOPROTEIN.

FT CHAIN 19 609

COPPER AND NICKEL (BY SIMILARITY).

FT METAL 22 22

BY SIMILARITY.

FT DISULFID 99 114

BY SIMILARITY.

FT DISULFID 113 134

BY SIMILARITY.

FT DISULFID 148 193

BY SIMILARITY.

FT DISULFID 192 201

BY SIMILARITY.

FT DISULFID 224 270

BY SIMILARITY.

FT DISULFID 269 277

BY SIMILARITY.

FT DISULFID 289 303

BY SIMILARITY.

FT DISULFID 302 313

BY SIMILARITY.

FT DISULFID 384 393

BY SIMILARITY.

FT DISULFID 416 452

BY SIMILARITY.

FT DISULFID 461 472

BY SIMILARITY.

FT DISULFID 485 501

BY SIMILARITY.

FT DISULFID 500 511

BY SIMILARITY.

FT DISULFID 538 583

BY SIMILARITY.

FT DISULFID 582 591

BY SIMILARITY.

FT CARBOHYD 251 251

N-LINKED (GLCNAC. . .) (POTENTIAL).

FT SEQUENCE 609 AA; 68697 MW; E8AE548377DB60EB CRC64;

Query Match 69.6%; Score 32; DB 1; Length 609;
Best Local Similarity 50.0%; Pred. No. 1e+02;
Matches 4; Conservative 4; Mismatches 0; Indels 0; Gaps 0;

Qy 1 QKLCQK 8

:|:|:|:

Db 96 EELCHERE 103

RESULT 11

FETA_HUMAN

ID FETA_HUMAN STANDARD; PRT; 609 AA.

AC P02771;

DT 21-JUL-1986 (Rel. 01, Created)

DT 21-JUL-1986 (Rel. 01, Last sequence update)

DT 20-AUG-2001 (Rel. 40, Last annotation update)

DE ALPHA-FETOPROTEIN PRECURSOR (ALPHA-FETOglobulin) (ALPHA-1-

DE FETOPROTEIN).

GN APP.

OS Homo sapiens (Human).

OC Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi;

OC Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.

OX NCBI_TaxID=9606;

RN [1]

RP SEQUENCE FROM N.A.

RX MEDLINE=6373664; PubMed=6192439;

RA Morinaga T., Sakai M., Wegmann T.G., Tamaoki T.;

RT "Primary structures of human alpha-fetoprotein and its mRNA.";

RL Proc. Natl. Acad. Sci. U.S.A. 80:4604-4608(1983).

RN [2]

RP SEQUENCE FROM N.A.

RX MEDLINE=87185438; PubMed=2436661;

RA Gibbs P.E.M., Zielinski R., Boyd C., Dugaiczak A.;

RT "Structure, polymorphism, and novel repeated DNA elements revealed by

RT a complete sequence of the human alpha-fetoprotein gene.";

RL Biochemistry 26:1332-1343(1987).

RN SEQUENCE OF 1-28 FROM N.A.
RX MEDLINE=93278385; PubMed=7694942;
RA McVey J.H., Michaelides K., Hansen L.P., Ferguson-Smith M.,
RA Tilghman S., Krumlauf R., Tuddenham E.G.D.;

RT "A G-->A substitution in an HNF I binding site in the human alpha-fetoprotein gene is associated with hereditary persistence of alpha-fetoprotein (HFAFP).";
RL Hum. Mol. Genet. 2:379-379(1993).

[4]
RN SEQUENCE OF 429-556 FROM N.A.
RX MEDLINE=83158778; PubMed=6187626;
RA Beattie W.G., Dugaiczak A.;

RT "Structure and evolution of human alpha-fetoprotein deduced from partial sequence of cloned cDNA.";

RL Gene 20:415-422(1982).

[5]
RN PARTIAL SEQUENCE OF 19-609.
RX MEDLINE=91242409; PubMed=1709810;
RA Pucci P., Siciliano R., Malorni A., Marino G., Tecce M.F.,
RA Cecarini C., Terrana B.;

RT "Human alpha-fetoprotein primary structure: a mass spectrometric study.";

RL Biochemistry 30:5061-5066(1991).

[6]
RN PRELIMINARY SEQUENCE OF 19-35.
RX MEDLINE=77242506; PubMed=70228;
RA Yachnin S., Hsu R., Heinrikson R.L., Miller J.B.;

RT "Studies on human alpha-fetoprotein. Isolation and characterization of monomeric and polymeric forms and amino-terminal sequence analysis.";

RL Biochim. Biophys. Acta 493:418-428(1977).

[7]
RN PRELIMINARY SEQUENCE OF 19-38.
RX MEDLINE=78001760; PubMed=71198;
RA Aoyagi Y., Ikenaka T., Ichida F.;

RT "Comparative chemical structures of human alpha-fetoproteins from fetal serum and from ascites fluid of a patient with hepatoma.";

RL Cancer Res. 37:3663-3667(1977).

[8]
RN PRELIMINARY SEQUENCE OF 19-39.
RX MEDLINE=75018719; PubMed=4138095;
RA Ruoslahti E., Pihko H., Vaheeri A., Seppala M., Virolainen M.,
RA Kontinen A.;

RT "Alpha fetoprotein: structure and expression in man and inbred mouse strains under normal conditions and liver injury.";

RL Johns Hopkins Med. J. Suppl. 3:249-255(1974).

[9]
RN GENE STRUCTURE.
RX MEDLINE=85182629; PubMed=2580830;
RA Sakai M., Morinaga T., Urano Y., Watanabe K., Wegmann T.G.,
RA Tamaoki T.;

RT "The human alpha-fetoprotein gene. Sequence organization and the 5' flanking region.";

RL J. Biol. Chem. 260:5055-5060(1985).

[10]
RN METAL-BINDING.
RX MEDLINE=79001617; PubMed=80265;
RA Aoyagi Y., Ikenaka T., Ichida F.;

RT "Copper(II)-binding ability of human alpha-fetoprotein.";

RL Cancer Res. 38:3483-3486(1978).

[11]
RN BILIRUBIN-BINDING.
RX MEDLINE=80001710; PubMed=89900;
RA Aoyagi Y., Ikenaka T., Ichida F.;

RT "Alpha-fetoprotein as a carrier protein in plasma and its bilirubin-binding ability.";

RL Cancer Res. 39:3571-3574(1979).

[12]
RN SULFATION.
RX PubMed=2414772;
RA Liu M.C., Yu S., Sy J., Redman C.M., Lipmann F.;

RT "Tyrosine sulfation of proteins from the human hepatoma cell line

RT RepG2.":
RL Proc. Natl. Acad. Sci. U.S.A. 82:7160-7164(1985).
CC -!- FUNCTION: BINDS COPPER, NICKEL, AND FATTY ACIDS AS WELL AS, AND
CC BILIRUBIN LESS WELL THAN, SERUM ALBUMIN. ONLY A SMALL PERCENTAGE
CC (LESS THAN 2%) OF THE HUMAN AFP SHOWS ESTROGEN-BINDING PROPERTIES.
CC -!- SUBUNIT: DIMERIC AND TRIMERIC FORMS HAVE BEEN FOUND IN ADDITION
CC TO THE MONOMERIC FORM.
CC -!- SUBCELLULAR LOCATION: EXTRACELLULAR.
CC -!- TISSUE SPECIFICITY: PLASMA. SYNTHESIZED BY THE FETAL LIVER AND
CC YOLK SAC.
CC -!- DEVELOPMENTAL STAGE: OCCURS IN THE PLASMA OF FETUSES MORE THAN 4
CC WEEKS OLD, REACHES THE HIGHEST LEVELS DURING THE 12TH-16TH WEEK OF
CC GESTATION, AND DROPS TO TRACE AMOUNTS AFTER BIRTH. THE SERUM LEVEL
CC IN ADULTS IS USUALLY LESS THAN 40 NG/ML. AFP OCCURS ALSO AT HIGH
CC LEVELS IN THE PLASMA AND ASCITIC FLUID OF ADULTS WITH HEPATOMA.
CC -!- DOMAIN: COMPOSED OF THREE HOMOLOGOUS DOMAINS.
CC -!- PTM: INDEPENDENT STUDIES SUGGEST HETEROGENEITY OF THE AMINO-
CC TERMINAL SEQUENCE OF THE MATURE PROTEIN AND OF THE CLEAVAGE SITE
CC OF THE SIGNAL SEQUENCE.
CC -!- PTM: SULFATED.
CC -!- SIMILARITY: BELONGS TO THE ALB/AFP/VDB FAMILY.
CC -----
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CC -----
DR EMBL; M10949; AAA51674.1; -
DR EMBL; M10950; AAA51675.1; -
DR EMBL; V01514; CAA24759.1; -
DR EMBL; M16110; AAB58754.1; -
DR EMBL; J00077; AAC95396.1; -
DR EMBL; Z19532; CAA79592.1; -
DR PIR; A03234; FPHU
DR PIR; A26624; A26624.
DR HSP; P02768; IUOR.
DR GlycoSuiteDB; P02771; -
DR MIM; 104150; -
DR InterPro; IPR000264; Serum_albumin.
DR Pfam; PF00273; transport_prot; 3.
DR PRINTS; PR00802; SERUMALBUMIN.
DR PRINTS; PR00803; AFPETOPROTEIN.
DR SMART; SM00103; ALBUMIN; 3.
DR PROSITE; PS00212; ALBUMIN; 2.
KW Glycoprotein; Sulfation; Albumin; Plasma; Embryo; Repeat;
KW Metal-binding; Copper; Nickel; Signal.
FT SIGNAL 1 18
FT CHAIN 19 609 ALPHA-FETOPROTEIN.
FT METAL 22 22 COPPER AND NICKEL.
FT DISULFID 99 114
FT DISULFID 113 124
FT DISULFID 148 193
FT DISULFID 192 201
FT DISULFID 224 270
FT DISULFID 269 277
FT DISULFID 289 303
FT DISULFID 302 313
FT DISULFID 384 393
FT DISULFID 416 462
FT DISULFID 461 472
FT DISULFID 485 501
FT DISULFID 500 511
FT DISULFID 538 583
FT DISULFID 582 591
FT CARBOHYD 251 251 N-LINKED (GLCNAC...).
FT FTID=CAR_000070.
SEQUENCE 609 AA: 68677 MW: 4045820E1C2D4F CRC64;

Query Match 69.6%; Score 32; DB 1; Length 609;

Best Local Similarity 50.0%; Pred. No. 1e+02; Matches 4; Conservative 4; Mismatches 0; Indels 0; Gaps 0;
QY 1 QKLCQK 8
DB 96 BELCHEKE 103
RESULT 12
Y081_CAEEL STANDARD; PRT; 645 AA.
AC P34617;
DT 01-FEB-1994 (Rel. 28, Created)
DT 01-FEB-1995 (Rel. 31, Last sequence update)
DT 20-AUG-2001 (Rel. 40, Last annotation update)
DE HYPOTHETICAL GTP-BINDING PROTEIN ZK1236.1 IN CHROMOSOME III.
GN ZK1236.1.
OS Caenorhabditis elegans.
OC Eukaryota; Metazoa; Nematoda; Chromadorea; Rhabditida; Rhabditoidea;
OC Rhabditidae; Peloderinae; Caenorhabditis.
OX NCBI_TaxID=6239;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=BRISTOL N2;
RX MEDLINE=94150718; PubMed=7906398;
RA Wilson R., Ainscough R., Anderson K., Baynes C., Berks M., Coulson A.,
RA Bonfield J., Burton J., Connell M., Copsey T., Cooper J., Fraser A.,
RA Craxton M., Dear S., Du Z., Durbin R., Favello A., Fraser A.,
RA Fulton L., Gardner A., Green P., Hawkins T., Hillier L., Jier M.,
RA Johnston L., Jones M., Kershaw J., Kirsten J., Laister N.,
RA Latreille P., Lightning J., Lloyd C., Mortimore B., O'Callaghan M.,
RA Parsons J., Percy C., Rifken L., Roopra A., Saunders D., Shownkeen R.,
RA Sims M., Smailson N., Smith A., Smith M., Sonhammer E., Staden R.,
RA Sulston J., Thierry-Mieg J., Thomas K., Vaudin M., Vaughan K.,
RA Watson R., Watson A., Weinstock L., Wilkinson-Sproat J.,
RA Wohldman P.;
RT "2.2 Mb of contiguous nucleotide sequence from chromosome III of C.
RT elegans.";
RL Nature 368:32-38(1994).
CC -!- SIMILARITY: BELONGS TO THE GTP-BINDING ELONGATION FACTOR FAMILY.
CC LEPA SUBFAMILY.
CC -----
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CC -----
DR EMBL; L13200; AAA28191.2; -
DR HSP; P13551; IELO.
DR WormRep; ZK1236.1; CE01446.
DR InterPro; IPR000795; GTP_EFTU.
DR Pfam; PF00009; GTP_EFTU; 1.
DR PROSITE; PS00301; EFATOR_GTP; 1.
KW Hypothetical protein; GTP-binding.
FT NP_BIND 49 56 GTP (POTENTIAL).
FT NP_BIND 108 112 GTP (POTENTIAL).
FT NP_BIND 162 165 GTP (POTENTIAL).
FT SEQUENCE 645 AA: 72268 MW: 3F08EA3E5FD53819 CRC64;
Query Match 69.6%; Score 32; DB 1; Length 645;
Best Local Similarity 75.0%; Pred. No. 1.1e+02; Matches 6; Conservative 1; Mismatches 1; Indels 0; Gaps 0;
QY 1 QKLCQK 8
DB 614 KKLHQK 621
RESULT 13

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SVI_MYCGE
ID SYI_MYCGE STANDARD; PRT; 895 AA.
AC P47587;
DT 01-FEB-1996 (Rel. 33, Created)
DT 01-FEB-1996 (Rel. 33, Last sequence update)
DT 20-AUG-2001 (Rel. 40, Last annotation update)
DE ISOLEUCYL-TRNA SYNTHETASE (EC 6.1.1.5) (ISOLEUCINE--TRNA LIGASE)
DE (ILERS).
DE
GN ILRS OR MG345.
OS Mycoplasma genitalium.
OC Bacteria; Firmicutes; Bacillus/Clostridium group; Mollicutes;
OC Mycoplasmataceae; Mycoplasma.
OX NCBI_TaxID=2097;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=ATCC 33530 / G-37;
RX MEDLINE=96026346; PubMed=7569993;
RA Fraser C.M., Gocayne J.D., White O., Adams M.D., Clayton R.A.,
RA Fleischmann R.D., Bult C.J., Kerlavage A.R., Sutton G., Kelley J.M.,
RA Fritchman J.L., Weidman J.F., Small K.V., Sandusky M., Fuhrmann J.L.,
RA Nguyen D.T., Utterback T.R., Saudek D.M., Phillips C.A., Merrick J.M.,
RA Tomb J.-F., Dougherty B.A., Boff K.F., Hu P.-C., Lucier T.S.,
RA Peterson S.N., Smith H.O., Hutchison C.A. III, Venter J.C.;
RT "The minimal gene complement of Mycoplasma genitalium.";
RL Science 270:397-403(1995).
RN [2]
RP SEQUENCE OF 262-371 AND 605-711 FROM N.A.
RC STRAIN=ATCC 33530 / G-37;
RX MEDLINE=94075230; PubMed=8253680;
RA Peterson S.N., Hu P.-C., Boff K.F., Hutchison C.A. III;
RT "A survey of the Mycoplasma genitalium genome by using random
RT sequencing.";
RL J. Bacteriol. 175:7918-7930(1993).
CC -!- CATALYTIC ACTIVITY: ATP + L-ISOLEUCINE + TRNA(ILE) = AMP +
CC PYROPHOSPHATE + L-ISOLEUCYL-TRNA(ILE).
CC -!- COPACATOR: BINDS ONE ZINC ION (BY SIMILARITY).
CC -!- SUBUNIT: MONOMER (BY SIMILARITY).
CC -!- SUBCELLULAR LOCATION: CYTOPLASMIC.
CC -!- SIMILARITY: BELONGS TO CLASS-I AMINOACYL-TRNA SYNTHETASE FAMILY.
CC
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CC or send an email to license@isb-sib.ch).
CC
CC EMBL; D84432; BAA12574.1; ALT_INIT.
CC EMBL; 299116; CAB4360.1; ALT_INIT.
CC Subtilist; BG11713; xseB.
CC InterPro; IPR003761; Exonuc_VII_S.
CC Pfam; PF02609; Exonuc_VII_S; 1.
CC Hydrolase; Nuclease; Exonuclease; Complete proteome.
CC SEQUENCE 75 AA; 8571 MW; 6A28593DAC9F092A CRC64;
CC
CC Query Match 67.4%; Score 31; DB 1; Length 75;
CC Best Local Similarity 66.7%; Pred. No. 25;
CC Matches 4; Conservative 2; Mismatches 0; Indels 0; Gaps 0;
CC
CC QY 2 KLCHQK 7
CC I:||||
CC Db 39 KMCHEK 44
CC
CC RESULT 15
CC NEUT_BOVIN
CC ID NEUT_BOVIN STANDARD; PRT; 169 AA.
CC AC P01156;
CC DT 21-JUL-1986 (Rel. 01, Created)
CC DT 01-JUL-1989 (Rel. 11, Last sequence update)
CC DT 15-DEC-1998 (Rel. 37, Last annotation update)
CC DE NEUROTENSIN/NEUROMEDIN N PRECURSOR (NT/NMN).
CC NTS
CC Bos taurus (Bovine).
CC OS Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
CC Mammalia; Eutheria; Cetartiodactyla; Ruminantia; Pecora; Bovidae;
CC Bovidae; Bovinae; Bos.
CC OX NCBI_TaxID=9913;
CC RN [1]

```



```

RP SEQUENCE FROM N.A.
RX MEDLINE=88169625; PubMed=2832414;
RA Kislaukis E., Bullock B., McNeil S., Dobner P.R.;
RT "The rat gene encoding neurotensin and neuromedin N. Structure,
RT tissue-specific expression, and evolution of exon sequences.";
RL J. Biol. Chem. 263:4963-4968(1988).
RN [2]
RP SEQUENCE OF 150-162.
RC TISSUE=Hypothalamus;
RX MEDLINE=75095678; PubMed=1167549;
RA Caraway R., Leeman S.E.;
RT "The amino acid sequence of a hypothalamic peptide, neurotensin.";
RL J. Biol. Chem. 250:1907-1911(1975).
RN [3]
RP SYNTHESIS OF NEUROTENSIN.
RX MEDLINE=75095679; PubMed=1112838;
RA Caraway R., Leeman S.E.;
RT "The synthesis of neurotensin.";
RL J. Biol. Chem. 250:1912-1918(1975).
CC -1- FUNCTION: NEUROTENSIN MAY PLAY AN ENDOCRINE OR PARACRINE ROLE
CC IN THE REGULATION OF FAT METABOLISM. IT CAUSES CONTRACTION OF
CC SMOOTH MUSCLE.
CC -1- SUBCELLULAR LOCATION: PACKAGED WITHIN SECRETORY VESICLES.
CC -1- TISSUE SPECIFICITY: BRAIN AND GUT.
CC -1- SIMILARITY: C-TERMINAL SEQUENCE SIMILARITY WITH NEUROTENSIN-
CC RELATED PEPTIDES.
CC -----
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CC or send an email to license@isb-sib.ch).
CC -----
DR EMBL: M18621; AAA30668.1; -.
DR PIR: A01420; UNEO.
KW Cleavage on pair of basic residues; Vasoactive; Signal.
FT SIGNAL 1 22
FT CHAIN 23 147 LARGE NEUROMEDIN N (NMN-125).
FT PEPTIDE 142 147 NEUROMEDIN N.
FT PEPTIDE 150 162 NEUROTENSIN.
FT MOD_RES 150 150 PYRROLIDONE CARBOXYLIC ACID.
SQ SEQUENCE 169 AA; 19712 MW; 7B78760D5E4D7D32 CRC64;

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Query Match 67.4%; Score 31; DB 1; Length 169;
Best Local Similarity 57.1%; Pred. No. 51;
Matches 4; Conservative 2; Mismatches 1; Indels 0; Gaps 0;

Qy 1 QKLCQK 7
Db 107 QKICHSR 113

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Search completed: February 12, 2002, 12:04:02
Job time: 796 sec

GenCore version 4.5
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OM protein - protein search, using sw model

Run on: February 12, 2002, 11:51:40 ; Search time 55.4 Seconds
(without alignments)
11.000 Million cell updates/sec

Title: US-09-606-129A-19
Perfect score: 46
Sequence: 1 QKLCQKK 8
Scoring table: BLOSUM62
Gap 10.0 , Gapext 0.5

Searched: 219241 seqs, 76174552 residues
Total number of hits satisfying chosen parameters: 219241

Minimum DB seq length: 0
Maximum DB seq length: 2000000000
Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database : PIR_68:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match %	Length	DB ID	Description
1	46	100.0	295	2 A42268	billiverdin reductase
2	36	78.3	654	1 BVBYPI	CBP1 protein - yea
3	34	73.9	547	2 T39478	zinc-finger protei
4	34	73.9	1188	2 A71621	protein with 5'-3'
5	34	73.9	1295	2 T30528	reverse transcript
6	34	73.9	1473	2 T31422	C-terminal domain
7	33	71.7	263	2 E72675	hypothetical prote
8	33	71.7	673	2 S36336	probable transcrip
9	33	71.7	773	2 T00554	hypothetical prote
10	33	71.7	1247	2 T18671	hypothetical prote
11	32	69.6	135	1 W6WLEP	E6 protein - Europ
12	32	69.6	210	2 T16125	hypothetical prote
13	32	69.6	290	2 T39522	3-dehydroquinat
14	32	69.6	345	1 S26707	transcription init
15	32	69.6	357	2 E64233	membrane transport
16	32	69.6	380	2 T24081	hypothetical prote
17	32	69.6	422	2 E96753	hypothetical prote
18	32	69.6	462	2 A46170	tektin A1 - sea ur
19	32	69.6	465	2 T27032	hypothetical prote
20	32	69.6	474	2 T00699	hypothetical prote
21	32	69.6	548	2 T22337	hypothetical prote
22	32	69.6	581	2 S44896	ZK1236.1 protein -
23	32	69.6	595	2 F71471	probable DNA prima
24	32	69.6	609	1 PPHU	alpha-fetoprotein
25	32	69.6	609	1 PPGU	alpha-fetoprotein
26	32	69.6	895	2 B64238	isoleucine--trna l
27	32	69.6	920	2 JC7313	aryl hydrazocarbon
28	32	69.6	1056	2 T00060	hypothetical prote
29	31	67.4	54	2 H69960	exodeoxyribonuclea

30 31 67.4 170 1 UNDG
31 31 67.4 251 2 G86368
32 31 67.4 266 2 T34411
33 31 67.4 271 2 T16421
34 31 67.4 299 2 A75401
35 31 67.4 364 2 T03892
36 31 67.4 436 2 D84782
37 31 67.4 478 2 C71523
38 31 67.4 491 2 G83850
39 31 67.4 498 2 T11683
40 31 67.4 510 2 D96782
41 31 67.4 631 2 JC5803
42 31 67.4 670 2 B70145
43 31 67.4 676 2 T39766
44 31 67.4 704 2 T05565
45 31 67.4 977 2 S53302

neurotensin precur
hypothetical prote
hypothetical prote
hypothetical prote
ribulose-phosphate
hypothetical prote
probable prolina t
probable shikimate
Atp-dependent DNA
probable iron-sulf
hypothetical prote
ring finger protei
periplasmic protei
probable long-chain
hypothetical prote
H+-transporting At

ALIGNMENTS

RESULT 1

A42268
billiverdin reductase (EC 1.3.1.24) - rat
C:Species: Rattus norvegicus (Norway rat)
C:Date: 04-Mar-1993 #sequence_revision 18-Nov-1994 #text_change 05-Nov-1999
C:Accession: A42268
R:Fakhrat, H.; Maines, M.D.
J. Biol. Chem. 267, 4023-4029, 1992
A:Title: Expression and characterization of a cDNA for rat kidney billiverdin reductase
A:Reference number: A42268; MUID:92156147
A:Accession: A42268
A:Status: preliminary; not compared with conceptual translation
A:Molecule type: nucleic acid; protein
A:Residues: 1-295 <FA>
A:Cross-references: GB:M81681; NID:9203177; PIDN:AAA40830.1; PID:g203178
A:Experimental source: kidney
A:Note: sequence extracted from NCBI backbone (NCBIP:82800)
C:Keywords: liver; oxidoreductase

Query Match 100.0%; Score 46; DB 2; Length 295;
Best Local Similarity 100.0%; Pred. No. 0.37; Indels 0; Gaps 0;
Matches 8; Conservative 0; Mismatches 0;

QY 1 QKLCQKK 8
|||
DB 288 QKLCQKK 295

RESULT 2

BVBYPI
CBP1 protein - yeast (Saccharomyces cerevisiae)
N:Alternate names: protein HRA654; protein J0242; protein YJL209W
C:Species: Saccharomyces cerevisiae
C:Date: 31-Mar-1991 #sequence_revision 31-Mar-1991 #text_change 12-Nov-1999
C:Accession: S05829; S50776; S56999; S45164
R:Diekmann, C.L.; Homison, G.; Tzagoloff, A.
J. Biol. Chem. 259, 4732-4738, 1984
A:Title: Assembly of the mitochondrial membrane system. Nucleotide sequence of a y
A:Reference number: S05829; MUID:84185566
A:Accession: S05829
A:Molecule type: DNA
A:Residues: 1-654 <DIE>
A:Cross-references: EMBL:K02647; NID:g171166; PIDN:AAA34474.1; PID:g171167
R:Vandenbol, M.; Durand, P.; Bolle, P.A.; Dion, C.; Portetelle, D.; Hilger, F.
Yeast 10, 1657-1662, 1994
A:Title: Sequence analysis of a 40.2 kb DNA fragment located near the left telomer.
A:Reference number: S50701; MUID:95242842
A:Accession: S50776
A:Status: nucleic acid sequence not shown; translation not shown
A:Molecule type: DNA
A:Residues: 1-654 <YAW>

C;Cross-references: EMBL:Z34098; NID:g496934; PIDN:CAA84002.1; PID:g496953
 A;Note: the nucleotide sequence was submitted to the EMBL Data Library, June 1994
 R;Vandenbol, M.; Durand, P.; Portetelie, D.; Hilger, F.
 submitted to the Protein Sequence Database, September 1995
 A;Reference number: S56835
 A;Accession: S56999
 A;Molecule type: DNA
 A;Residues: 1-654 <VAN>
 C;Cross-references: EMBL:Z49484; NID:g1015590; PIDN:CAA89506.1; PID:g1015591; GSPDB:GN000100
 R;Purnelle, B.; Coster, F.; Goffeau, A.
 submitted to the Protein Sequence Database, September 1995
 A;Reference number: S56977
 A;Accession: S56996
 A;Molecule type: DNA
 A;Residues: 637-654 <PUR>
 C;Cross-references: EMBL:Z49484; GSPDB:GN00010; MIPS:YJL209W
 C;Genetics:
 A;Gene: SGD.CBP1; MIPS:YJL209W
 A;Cross-references: SGD:S0003745; MIPS:YJL209W
 A;Map position: 10L
 A;Genome: nuclear
 C;Function:
 A;Description: pre-mRNA processing
 A;Note: required for correct 5' terminal processing of cytochrome b pre-mRNA
 C;Superfamily: CBP1 protein
 C;Keywords: mitochondrion

Query Match 78.3%; Score 36; DB 1; Length 654;
 Best Local Similarity 75.0%; Pred. No. 44;
 Matches 6; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

Qy 1 QKLCCHK 8
 :|||||
 Db 514 KKLCCHK 521

RESULT 3

T39478
 zinc-finger protein - fission yeast (Schizosaccharomyces pombe)

C;Species: Schizosaccharomyces pombe
 C;Date: 03-Dec-1999 #sequence_revision 03-Dec-1999 #text_change 02-Sep-2000
 C;Accession: T39478
 R;Lyne, M.; Rajandream, M.A.; Barrell, B.G.; Lucas, M.; Gaillardin, C.
 submitted to the EMBL Data Library, August 1997

A;Reference number: Z21858
 A;Accession: T39478
 A;Status: preliminary; translated from GB/EMBL/DBBJ
 A;Molecule type: DNA
 A;Residues: 1-547 <LYN>

C;Cross-references: EMBL:AL031349; PIDN:CAA20477.1; GSPDB:GN00067; SPDB:SPBC15D4.02
 A;Experimental source: strain 972h-; cosmid c15D4
 C;Genetics:

A;Gene: SPDB:SPBC15D4.02

A;Map position: 2
 C;Superfamily: GAL4 zinc binuclear cluster homology
 F;169-206/Domain: GAL4 zinc binuclear cluster homology <GL4>

Query Match 73.9%; Score 34; DB 2; Length 547;
 Best Local Similarity 71.4%; Pred. No. 86;
 Matches 5; Conservative 2; Mismatches 0; Indels 0; Gaps 0;

Qy 1 QKLCCHK 7
 :|||||
 Db 427 RKLCHK 433

RESULT 4

A71621
 protein with 5'-3' exonuclease domain (Kem-1 family) PFB0205c - malaria parasite (Plasmodium falciparum)
 C;Species: Plasmodium falciparum
 C;Date: 13-Nov-1998 #sequence_revision 13-Nov-1998 #text_change 21-Jul-2000

C;Accession: A71621
 R;Gardner, M.J.; Tettelin, H.; Carucci, D.J.; Cummings, L.M.; Aravind, L.; Koonin, A.; Pertea, M.; Salzberg, S.; Zhou, L.; Sutton, G.G.; Clayton, R.; White, O.; Smith, S.
 Science 282, 1126-1132, 1998
 A;Title: Chromosome 2 sequence of the human malaria parasite Plasmodium falciparum.
 A;Reference number: A71600; MUID:99021743
 A;Accession: A71621
 A;Status: preliminary; nucleic acid sequence not shown; translation not shown
 A;Molecule type: DNA
 A;Residues: 1-1188 <GAR>
 A;Cross-references: GB:AE001380; GB:AE001362; NID:g3845120; PIDN:AA71830.1; PID:g3
 A;Experimental source: clone 3D7
 C;Genetics:
 A;Gene: PFB0205c

Query Match 73.9%; Score 34; DB 2; Length 1188;
 Best Local Similarity 71.4%; Pred. No. 1.6e+02;
 Matches 5; Conservative 2; Mismatches 0; Indels 0; Gaps 0;

Qy 1 QKLCCHK 7
 :|||||
 Db 1117 EELCHK 1123

RESULT 5

T30528
 reverse transcriptase - rice blast fungus

C;Species: Magnaporthe grisea (rice blast fungus)
 C;Date: 22-Oct-1999 #sequence_revision 22-Oct-1999 #text_change 22-Oct-1999
 C;Accession: T30528
 R;Meyn III, M.A.; Farrall, L.; Valent, B.; Chumley, F.G.; Orbach, M.J.
 submitted to the EMBL Data Library, August 1997

A;Description: Magnaporthe grisea repeated DNA element MGR583 is a member of the LI
 A;Reference number: Z20845
 A;Accession: T30528
 A;Status: preliminary; translated from GB/EMBL/DBBJ
 A;Molecule type: DNA
 A;Residues: 1-1295 <MEY>
 A;Cross-references: EMBL:AF018033; NID:g2454620; PID:g2454622; PIDN:AA71689.1

Query Match 73.9%; Score 34; DB 2; Length 1295;
 Best Local Similarity 71.4%; Pred. No. 1.7e+02;
 Matches 5; Conservative 2; Mismatches 0; Indels 0; Gaps 0;

Qy 2 KLCHKOK 8
 :|||||
 Db 33 ELCHKOK 39

RESULT 6

T31422
 C-terminal domain-binding protein RA9 - rat

C;Species: Rattus norvegicus (Norway rat)
 C;Date: 29-Oct-1999 #sequence_revision 29-Oct-1999 #text_change 07-Dec-1999
 C;Accession: T31422
 R;Ryryev, A.; Patturajan, M.; Litingtung, Y.; Joshi, R.V.; Gentile, C.; Gebata, M.; Proc. Natl. Acad. Sci. U.S.A. 93, 6975-6980, 1996
 A;Title: The C-terminal domain of the largest subunit of RNA polymerase II interact
 A;Reference number: Z21024; MUID:96293459

A;Accession: T31422
 A;Status: preliminary; translated from GB/EMBL/DBBJ
 A;Molecule type: mRNA

A;Residues: 1-1473 <YUR>
 A;Cross-references: EMBL:U49057; NID:g1438533; PID:g1438534; PIDN:AA52658.1
 A;Experimental source: hippocampus

Query Match 73.9%; Score 34; DB 2; Length 1473;
 Best Local Similarity 71.4%; Pred. No. 1.9e+02;
 Matches 5; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

QY 1 QKLCQK 7
11:1111
Db 1425 QKICHSK 1431

RESULT 7
E72675
hypothetical protein APE0826 - Aeropyrum pernix (strain K1)
C:Species: Aeropyrum pernix
C:Date: 20-Aug-1999 #sequence_revision 20-Aug-1999 #text_change 20-Jun-2000
R:Accession: E72675
R:Kawarayashi, Y.; Hino, Y.; Horikawa, H.; Yamazaki, S.; Haikawa, Y.; Jin-no, K.; Takahawa, H.; Takamiya, M.; Masuda, S.; Funahashi, T.; Tanaka, T.; Kudoh, Y.; Yamazaki, J.; K DNA Res. 6, 83-101, 1999
A:Title: Complete genome sequence of an aerobic hyper-thermophilic Crenarchaeon, Aeropyrum pernix strain K1
A:Reference number: A72450; MUID:99310339
A:Accession: E72675
A:Status: preliminary
A:Molecule type: DNA
A:Residues: 1-263 <RAW>
A:Cross-references: DDBJ:AF000060; NID:g5104188; PIDN:BAA79805.1; PID:g5104490
A:Experimental source: strain K1
C:Genetics:
A:Gene: APE0826
C:Superfamily: Aeropyrum pernix hypothetical protein APE0826

Query Match 71.7%; Score 33; DB 2; Length 263;
Best Local Similarity 83.3%; Pred. No. 73;
Matches 5; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

QY 3 LCHQK 8
111111
Db 258 LCHQK 263

RESULT 8
S36336
probable transcription factor PLZF - human
C:Species: Homo sapiens (man)
C:Date: 31-Dec-1993 #sequence_revision 31-Dec-1993 #text_change 05-Nov-1999.
R:Accession: S36336; S31989
R:Chen, Z.; Brand, N.J.; Chen, A.; Chen, S.J.; Tong, J.H.; Wang, Z.Y.; Waxman, S.; Zelen EMBO J. 12, 1161-1167, 1993
A:Title: Fusion between a novel Krueppel-like zinc finger gene and the retinoic acid receptor gene
A:Reference number: S36336; MUID:93209216
A:Accession: S36336
A:Molecule type: mRNA
A:Residues: 1-673 <CHE>
A:Cross-references: EMBL:Z19002; NID:g38517; PIDN:CAA79489.1; PID:g38518
C:Genetics:
A:Gene: PLZF
C:Superfamily: POZ domain homology
C:Keywords: zinc finger
F;20-118/Domain: POZ domain homology <POZ>

Query Match 71.7%; Score 33; DB 2; Length 673;
Best Local Similarity 83.3%; Pred. No. 1.5e+02;
Matches 5; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

QY 2 KLCHQK 7
111111
Db 605 KLCHQK 610

RESULT 9
T00554
hypothetical protein A2g99440 [imported] - Arabidopsis thaliana
N:Alternate names: hypothetical protein F12L6.10
C:Species: Arabidopsis thaliana (mouse-ear cress)
C:Date: 01-Feb-1999 #sequence_revision 01-Feb-1999 #text_change 16-Feb-2001
C:Accession: T00554; D84817

R:Rounsley, S.D.; Lin, X.; Ketchum, K.A.; Crosby, M.L.; Brandon, R.C.; Sykes, S.M.; submitted to the EMBL Data Library, July 1998
A:Description: Arabidopsis thaliana chromosome II BAC F12L6 genomic sequence.
A:Reference number: Z14168
A:Accession: T00554
A:Status: translated from GB/EMBL/DDBJ
A:Molecule type: DNA
A:Residues: 1-773 <ROU>
A:Cross-references: EMBL:AC004218; NID:g3355463; PID:g3355473
A:Experimental source: cultivar Columbia
R:Lin, X.; Kaul, S.; Rounsley, S.D.; Shea, T.P.; Benito, M.I.; Town, C.D.; Fujii, C.M.; Koo, H.; Moffat, K.S.; Cronin, L.A.; Shen, M.; Vanaken, S.E.; Unayam, L.; Tallent, D.; Nierman, W.C.; White, O.; Eisen, J.A.; Salzberg, S.L.; Fraser, C.M.; Venter Nature 402, 761-768, 1999
A:Title: Sequence and analysis of chromosome 2 of the plant Arabidopsis thaliana.
A:Reference number: A84420; MUID:20083487
A:Accession: D84817
A:Status: preliminary
A:Molecule type: DNA
A:Residues: 1-773 <STO>
A:Cross-references: GB:AE002093; NID:g3355473; PIDN:AAC27835.1; GSPDB:GN00139
C:Genetics:
A:Gene: At2g39440; F12L6.10
A:Map position: 2
A:Introns: 35/1; 75/1; 117/1; 159/1; 222/3; 283/3; 294/1; 506/3; 567/1

Query Match 71.7%; Score 33; DB 2; Length 773;
Best Local Similarity 75.0%; Pred. No. 1.7e+02;
Matches 6; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

QY 1 QKLCQK 8
111111
Db 462 KKLCQK 469

RESULT 10
TI8671
hypothetical protein B0240.2 - Caenorhabditis elegans
C:Species: Caenorhabditis elegans
C:Date: 15-Oct-1999 #sequence_revision 15-Oct-1999 #text_change 21-Jan-2000
C:Accession: TI8671
R:White, S.
submitted to the EMBL Data Library, June 1996
A:Reference number: Z19004
A:Accession: TI8671
A:Status: preliminary; translated from GB/EMBL/DDBJ
A:Molecule type: DNA
A:Residues: 1-1247 <WIL>
A:Cross-references: EMBL:Z74026; PIDN:CAA98416.1; GSPDB:GN00023; CESP:B0240.2
A:Experimental source: clone B0240
C:Genetics:
A:Gene: CESP:B0240.2
A:Map position: 5
A:Introns: 29/2; 73/2; 108/2; 129/3; 202/2; 265/3; 401/3; 454/1; 466/3; 553/3; 594/3; 594/3; 594/3
C:Superfamily: Caenorhabditis elegans hypothetical protein B0240.2

Query Match 71.7%; Score 33; DB 2; Length 1247;
Best Local Similarity 71.4%; Pred. No. 2.5e+02;
Matches 5; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

QY 2 KLCHQK 8
111111
Db 265 KICQAK 271

RESULT 11
W6WLEP
E6 protein - European elk papillomavirus
C:Species: European elk papillomavirus
C:Date: 31-Mar-1989 #sequence_revision 31-Mar-1989 #text_change 11-May-2000
C:Accession: A29499; F9457; F94506

R:Ahola, H.; Bergman, P.; Stroem, A.C.; Moreno-Lopez, J.; Pettersson, U.
 Gene 50, 195-205, 1986
 A:Title: Organization and expression of the transforming region from the European elk papilloma virus
 A:Reference number: A91567; MUID:87219878
 A:Accession: A29499
 A:Molecule type: DNA
 A:Residues: 1-135 <AHO>
 A:Cross-references: GB:M15953; NID:g333025; PIDN:AAA66849.1; PID:g484015
 R:Eriksson, A.
 unpublished results 1987, cited by GenBank
 A:Reference number: A94457
 A:Accession: F94457
 A:Molecule type: DNA
 A:Residues: 1-135 <ERI>
 A:Cross-references: GB:M15953; NID:g333025; PIDN:AAA66849.1; PID:g484015
 R:Pettersson, U.
 submitted to GenBank, August 1987
 A:Reference number: A94506
 A:Accession: F94506
 A:Molecule type: DNA
 A:Residues: 1-135 <PET>
 A:Cross-references: GB:M15953; NID:g333025; PIDN:AAA66849.1; PID:g484015
 C:Superfamily: papillomavirus E6 protein
 C:Keywords: DNA binding; early protein; zinc finger

Query Match 69.6%; Score 32; DB 1; Length 135;
 Best Local Similarity 71.4%; Pred. No. 65;
 Matches 5; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

QY 2 KLCHQKK 8
 |||||
 Db 24 KRCHKK 30

RESULT 12
 T16125
 hypothetical protein F21H12.3 - Caenorhabditis elegans
 C:Species: Caenorhabditis elegans
 C:Date: 20-Sep-1999 #sequence_revision 20-Sep-1999 #text_change 24-Nov-1999
 C:Accession: T16125
 R:Favell, T.
 submitted to the EMBL Data Library, July 1995
 A:Description: The sequence of C. elegans cosmid F21H12.
 A:Reference number: Z18464
 A:Accession: T16125
 A:Status: preliminary; translated from GB/EMBL/DBJ
 A:Molecule type: DNA
 A:Residues: 1-210 <FAV>
 A:Cross-references: EMBL:U23176; NID:g726404; PID:g726407; PIDN:AAC46715.1; CESP:F21H12.
 A:Experimental source: strain Bristol N2
 C:Genetics:
 A:Gene: CESP:F21H12.3
 A:Introns: 37/3; 62/2
 C:Superfamily: Caenorhabditis elegans hypothetical protein F21H12.3

Query Match 69.6%; Score 32; DB 2; Length 210;
 Best Local Similarity 71.4%; Pred. No. 92;
 Matches 5; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

QY 1 QKCHQK 7
 :|||||
 Db 23 KNLCHQK 29

RESULT 13
 I39522
 3-dehydroquinase dehydratase (EC 4.2.1.10), catabolic [validated] - Acinetobacter calcoaceticus
 C:Species: Acinetobacter calcoaceticus
 C:Date: 19-Jul-1996 #sequence_revision 19-Jul-1996 #text_change 20-Jun-2000
 C:Accession: I39522
 R:Elsom, D.A.; Ornston, L.N.

J. Bacteriol. 177, 5971-5978, 1995
 A:Title: Unusual ancestry of dehydratases associated with quinate catabolism in *Acinetobacter calcoaceticus*
 A:Reference number: I39522; MUID:96011389
 A:Accession: I39522
 A:Status: preliminary; translated from GB/EMBL/DBJ
 A:Molecule type: DNA
 A:Residues: 1-290 <RES>
 A:Cross-references: EMBL:U20284; NID:g644872; PID:g644873
 C:Genetics:
 A:Gene: quib
 C:Function:
 A:Description: EC 4.2.1.10 [validated; MUID:96011389]
 C:Superfamily: 3-dehydroquinase dehydratase; 3-dehydroquinase dehydratase homology
 C:Keywords: carbon-oxygen lyase; hydro-lyase
 F:38-277/Domain: 3-dehydroquinase dehydratase homology <DQD>

Query Match 69.6%; Score 32; DB 2; Length 290;
 Best Local Similarity 85.7%; Pred. No. 1.4e+02;
 Matches 6; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 2 KLCHQKK 8
 |||||
 Db 167 KLAHQKK 173

RESULT 14
 S26707
 transcription initiation factor IIF - yeast (Saccharomyces cerevisiae)
 N:Alternate names: protein P9513.4; protein YPR086W
 C:Species: Saccharomyces cerevisiae
 C:Date: 10-Sep-1999 #sequence_revision 10-Sep-1999 #text_change 21-Jul-2000
 C:Accession: S26707; S69072
 R:Pinto, I.; Ware, D.E.; Hampsey, M.
 Cell 68, 977-988, 1992
 A:Title: The yeast SUA7 gene encodes a homolog of human transcription factor TFIIB
 A:Reference number: S26707; MUID:92191276
 A:Accession: S26707
 A:Molecule type: DNA
 A:Residues: 1-345 <PIN>
 A:Cross-references: EMBL:M81380; NID:gl72776; PIDN:AAA35126.1; PID:gl72777
 A:Experimental source: strain T15
 R:Couch, J.
 submitted to the EMBL Data Library, March 1996
 A:Description: The sequence of S. cerevisiae cosmid 9513.
 A:Reference number: S69057
 A:Accession: S69072
 A:Molecule type: DNA
 A:Residues: 1-345 <COU>
 A:Cross-references: EMBL:U51033; NID:gl230676; PIDN:AAB68135.1; PID:gl230692; GSPDB
 C:Genetics:
 A:Gene: SGD:SUA7; MIPS:YPR086W
 A:Cross-references: SGD:S0006290; MIPS:YPR086W
 A:Map position: 18R
 C:Superfamily: transcription initiation factor IIF; transcription initiation factor
 C:Keywords: DNA binding; duplication; nucleus; transcription initiation; zinc finger
 F:23-318/Domain: transcription initiation factor IIF homology <TF2B>
 F:24-48/Region: zinc finger CCCC motif
 F:133-210,239-313/Region: duplication

Query Match 69.6%; Score 32; DB 1; Length 345;
 Best Local Similarity 71.4%; Pred. No. 1.4e+02;
 Matches 5; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

QY 2 KLCHQKK 8
 |||||:|
 Db 155 KLCHDEK 161

RESULT 15
 E64233
 membrane transport protein (glnQ) homolog MG303 - Mycoplasma genitalium

C;Species: Mycoplasma genitalium
C;Date: 17-Nov-1995 #sequence_revision 17-Nov-1995 #text_change 02-Feb-2001
C;Accession: E64233
R;Fraser, C.M.; Gocayne, J.D.; White, O.; Adams, M.D.; Clayton, R.A.; Fleischmann, R.D.;
M.; Fuhrmann, J.; Nguyen, D.; Utterback, T.R.; Saudek, D.M.; Phillips, C.A.; Merrick, J.
C.A.; Venter, J.C.
Science 270, 397-403, 1995
A;Title: The minimal gene complement of Mycoplasma genitalium.
A;Reference number: A64200; MUID:96026346
A;Accession: E64233
A;Status: preliminary; nucleic acid sequence not shown; translation not shown
A;Molecule type: DNA
A;Residues: 1-357 <TIGR>
A;Cross-references: GB:U39711; GB:L43967; NID:gl045997; PID:gl046002; TIGR:MG303
A;Experimental source: strain G-37
C;Genetics:
A;Genetic code: SGC3
C;Superfamily: unassigned ATP-binding cassette proteins; ATP-binding cassette homology
C;Keywords: ATP; nucleotide binding; P-loop
F;90-288/Domain: ATP-binding cassette homology <ABC>
F;107-114/Region: nucleotide-binding motif A (P-loop)

Query Match 69.6%; Score 32; DB 2; Length 357;
Best Local Similarity 83.3%; Pred. No. 1.4e+02;
Matches 5; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

Qy 2 KLCHK 7
 |||||
Db 183 KLCHK 188

Search completed: February 12, 2002, 11:51:40
Job time: 301 sec

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OM protein - protein search, using sw model

Run on: February 12, 2002, 11:49:43 ; Search time 98.92 Seconds
(without alignments)
5.991 Million cell updates/sec

Title: US-09-606-129A-19
Perfect score: 46
Sequence: 1 QKLCQKK 8

Scoring table: BLOSUM62
Gapop 10.0 , Gapext 0.5

Searched: 522463 seqs, 74073290 residues

Total number of hits satisfying chosen parameters: 522463

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database : A_Geneseq_1101.*

1: /SID88/gcgdata/geneseq/geneseq/AA1980.DAT.*
2: /SID88/gcgdata/geneseq/geneseq/AA1981.DAT.*
3: /SID88/gcgdata/geneseq/geneseq/AA1982.DAT.*
4: /SID88/gcgdata/geneseq/geneseq/AA1983.DAT.*
5: /SID88/gcgdata/geneseq/geneseq/AA1984.DAT.*
6: /SID88/gcgdata/geneseq/geneseq/AA1985.DAT.*
7: /SID88/gcgdata/geneseq/geneseq/AA1986.DAT.*
8: /SID88/gcgdata/geneseq/geneseq/AA1987.DAT.*
9: /SID88/gcgdata/geneseq/geneseq/AA1988.DAT.*
10: /SID88/gcgdata/geneseq/geneseq/AA1989.DAT.*
11: /SID88/gcgdata/geneseq/geneseq/AA1990.DAT.*
12: /SID88/gcgdata/geneseq/geneseq/AA1991.DAT.*
13: /SID88/gcgdata/geneseq/geneseq/AA1992.DAT.*
14: /SID88/gcgdata/geneseq/geneseq/AA1993.DAT.*
15: /SID88/gcgdata/geneseq/geneseq/AA1994.DAT.*
16: /SID88/gcgdata/geneseq/geneseq/AA1995.DAT.*
17: /SID88/gcgdata/geneseq/geneseq/AA1996.DAT.*
18: /SID88/gcgdata/geneseq/geneseq/AA1997.DAT.*
19: /SID88/gcgdata/geneseq/geneseq/AA1998.DAT.*
20: /SID88/gcgdata/geneseq/geneseq/AA1999.DAT.*
21: /SID88/gcgdata/geneseq/geneseq/AA2000.DAT.*
22: /SID88/gcgdata/geneseq/geneseq/AA2001.DAT.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	36	78.3	91	22 AAB75580	Human secreted pro
2	35	76.1	551	20 AAY29955	Mouse CGICE protei
3	34	73.9	88	21 AAB56915	Human prostate can
4	34	73.9	102	22 AAG76544	Human colon cancer
5	34	73.9	522	22 AAB95513	Human protein sequ
6	34	73.9	790	22 AAB95045	Human protein sequ
7	34	73.9	1188	21 AAB18183	Plasmodium falcipa
8	33	71.7	197	21 AAG08675	Arabidopsis thalia
9	33	71.7	242	21 AAG08674	Arabidopsis thalia
10	33	71.7	678	22 AAM25801	Human protein sequ
11	33	71.7	778	21 AAY79180	Haematopoietic ste

12	69.6	23	19	AAW79501	Wild-type yeast tr
13	69.6	23	19	AAW79502	Loss-of-function m
14	69.6	23	19	AAW79503	Loss-of-function m
15	69.6	23	19	AAW79504	Loss-of-function m
16	69.6	23	19	AAW79505	Loss-of-function m
17	69.6	23	19	AAW79506	Loss-of-function m
18	69.6	23	19	AAW79507	Loss-of-function m
19	69.6	23	19	AAW79508	Loss-of-function m
20	69.6	23	19	AAW79509	Loss-of-function m
21	69.6	23	19	AAW79510	Loss-of-function m
22	69.6	55	22	AAW36177	Peptide #10214 enc
23	69.6	88	22	AAW15135	Peptide #1569 enco
24	69.6	88	22	AAW27590	Peptide #1627 enco
25	69.6	88	22	AAW02876	Peptide #1558 enco
26	69.6	111	20	AAY37346	Amino acid sequenc
27	69.6	196	11	AAR05285	Amino acid sequenc
28	69.6	197	17	AAR99228	Recombinant human
29	69.6	197	22	AAB62068	Mature rhuAFP doma
30	69.6	289	22	AAU14513	Human novel protei
31	69.6	389	17	AAR93222	Recombinant human
32	69.6	389	22	AAB62071	Mature rhuAFP doma
33	69.6	398	20	AAW78169	Human secreted pro
34	69.6	404	22	AAU14277	Human novel protei
35	69.6	590	17	AAW01023	Alpha-foetoprotein
36	69.6	609	17	AAR99227	Human alpha-foetop
37	69.6	609	22	AAU07130	Human Alpha-fetopro
38	69.6	609	22	AAB62067	Human alpha-fetopr
39	69.6	920	22	AAB97388	Dioxin receptor am
40	67.4	53	21	AAG02555	Human secreted pro
41	67.4	63	22	AAU14489	Human novel protei
42	67.4	71	22	AAU14253	Human novel protei
43	67.4	80	21	AAB53346	Human colon cancer
44	67.4	154	19	AAW37864	Human protein comp
45	67.4	174	21	AAW82680	Human granulocyte

ALIGNMENTS

RESULT 1
AAB75580
ID AAB75580 standard; Protein; 91 AA.
XX
XX AAB75580;
AC
DT 06-APR-2001 (first entry)
XX Human secreted protein sequence encoded by gene 28 SEQ ID NO:134.

Human; secreted protein; immunosuppressive; antiarthritic; antirheumatic;
antiproliferative; cytostatic; cardiant; vasotropic; cerebroprotective;
neotrophic; neuroprotective; antibacterial; virucide; fungicide;
ophthalmological; vulnery; autoimmune disease; cardiovascular disorder;
hyperproliferative disorders; cerebrovascular disorder; wound healing;
nervous system disorder; ocular disorder; skin aging; chemotaxis;
food additive.

XX Homo sapiens.
OS
XX
PN WO200077026-A1.
XX 21-DEC-2000.

XX 01-JUN-2000; 2000WO-US14973.
XX
PR 11-JUN-1999; 99US-0138630.
XX
XX (HUMA-) HUMAN GENOME SCI INC.
PA (ROSE/) ROSEN C A.
XX
XX Rosen CA, Ruben SM, Komatsoulis GA;
XX WPI; 2001-071258/08.
DR

DR N-PSDB; AAF64203.
 XX Nucleic acid molecules encoding human secreted proteins, used in
 PT preventing, treating or ameliorating a disorder, e.g. Alzheimer's and
 PT Parkinson's diseases and cancers
 PT
 XX Disclosure; Page 60; 542pp; English.
 XX
 CC Human secreted proteins AAB75506 - AAB75554 are encoded by polynucleotide
 CC sequences AAF64176 - AAF64224. The specification includes amino acid
 CC sequences AAB75555 - AAB75606 which represent fragments of the human
 CC secreted proteins, and protein sequences with which they share homology.
 CC The proteins and polynucleotides, their agonists and antagonists have
 CC activities dependent on the tissues and cells in which they are
 CC expressed, examples of these activities include, immunosuppressive;
 CC antiarthritic; antirheumatic; antiproliferative; cytostatic; cardiant;
 CC vasotropic; cerebroprotective; neotropic; neuroprotective; antibacterial;
 CC virucide; fungicide; ophthalmological; and vulnerary. The proteins,
 CC polynucleotides, agonists and antagonists can be used to treat or detect
 CC or diagnose various diseases and disorders including, autoimmune
 CC diseases e.g. rheumatoid arthritis, hyperproliferative disorders
 CC e.g. neoplasms of the breast or liver, cardiovascular disorders
 CC e.g. cardiac arrest, cerebrovascular disorders e.g. cerebral ischaemia,
 CC angiogenesis, nervous system disorders e.g. Alzheimer's disease,
 CC infections caused by bacteria, viruses and fungi and ocular disorders
 CC e.g. corneal infection. The polypeptides can also be used to aid wound
 CC healing and epithelial cell proliferation, to prevent skin ageing due to
 CC sunburn, to maintain organs before transplantation, for supporting cell
 CC culture of primary tissues, to regenerate tissues and in chemotaxis. The
 CC polypeptides can also be used as a food additive or preservative to
 CC increase or decrease storage capabilities. Included in the invention are
 CC polynucleotide sequences AAF64167 - AAF64175 and peptide AAB75505 which
 CC are used in the isolation, identification and characterisation of the
 CC proteins of the invention.
 XX
 XX Sequence 91 AA;
 SQ

Query Match 78.3%; Score 36; DB 22; Length 91;
 Best Local Similarity 100.0%; Pred. No. 20;
 Matches 6; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 QKLCQK 6
 Db 30 qkLcqh 35

RESULT 2
 AAY29955
 ID AAY29955 standard; Protein; 551 AA.
 AC AAY29955;
 XX
 XX
 DT 22-NOV-1999 (first entry)
 XX
 XX Mouse CG1CE protein sequence.
 DE
 XX
 XX CG1CE; Best's macular dystrophy; mutation; diagnosis; detection;
 KW BMD; age-related macular dystrophy.
 KW
 XX Mus sp.
 OS
 XX WC9943695-A1.
 PN
 XX
 PD 02-SEP-1999.
 XX
 XX 22-FEB-1999; 99WO-US03790.
 PF
 XX 25-FEB-1998; 98US-0075941.
 PR
 PR 18-DEC-1998; 98US-0112926.
 XX
 XX (MERI) MERCK & CO INC.
 PA (UYUP-) UNIV UPPSALA.
 PA

XX Petrukhin K, Caskey CT, Metzker M, Wadelius C;
 PI WPI; 1999-540560/45.
 XX N-PSDB; AA221229.
 DR
 XX Human and mouse polynucleotides encoding CG1CE polypeptides -
 PT
 XX Claim 7; Fig 8; 67pp; English.
 PS
 XX The present sequence represents the mouse CG1CE protein. When the CG1CE
 CC gene is mutated it is responsible for Best's macular dystrophy (BMD).
 CC Polynucleotides encoding CG1CE are useful for diagnosing whether a
 CC patient carries a mutation in the CG1CE gene. Normal and mutated
 CC CG1CE proteins are useful for identifying activators and/or inhibitors
 CC of these proteins, in order to treat BMD. The CG1CE gene offers a
 CC simpler and cheaper method of diagnosing BMD without the need for the
 CC presence of the patient. The gene may also be useful to discovering
 CC the genetic cause of age-related macular dystrophy.
 XX
 XX Sequence 551 AA;
 SQ

Query Match 76.1%; Score 35; DB 20; Length 551;
 Best Local Similarity 62.5%; Pred. No. 1.5e+02;
 Matches 5; Conservative 2; Mismatches 1; Indels 0; Gaps 0;

QY 1 QKLCQK 8
 Db 490 qeichmkk 497

RESULT 3
 AAB56915
 ID AAB56915 standard; Protein; 88 AA.
 AC AAB56915;
 XX
 XX 13-MAR-2001 (first entry)
 DT
 XX Human prostate cancer antigen protein sequence SEQ ID NO:1493.
 DE
 XX Human; prostate cancer; prostate cancer antigen; detection; diagnosis;
 KW neuroprotective; cytostatic; cardioactive; immunomodulatory; muscular;
 KW vulnary; gastrointestinal; nephrotropic; antiinfective; gynaecological;
 KW antibacterial; gene therapy; neural; immune; reproductive; renal;
 KW gastrointestinal; pulmonary; cardiovascular; proliferative disorder;
 KW wound; infectious disease.
 KW
 XX Homo sapiens.
 OS
 XX WO200055174-A1.
 PN
 XX 21-SEP-2000.
 PD
 XX 08-MAR-2000; 2000WO-US05988.
 PF
 XX 12-MAR-1999; 99US-0124270.
 PR
 XX (HUMA-) HUMAN GENOME SCI INC.
 PA (ROSE/) ROSEN C A.
 PA
 XX Rosen CA, Ruben SM;
 PI
 XX WPI; 2000-587513/55.
 DR N-PSDB; AAF16118.
 DR
 XX Prostate cancer associated gene sequences, referred to as prostate
 PT cancer antigens, useful for treatment, prevention, and diagnosis of
 PT disorders such as prostate cancer -
 XX
 XX Claim 11; Page 1930-1931; 2338pp; English.
 PS
 XX

CC AAF15566 to AAF16505 encode the human prostate cancer associated
 CC proteins, called prostate cancer antigens, given in AAB56363 to AAB57302.
 CC The prostate cancer antigens can have neuroprotective, cytostatic,
 CC cardioactive, immunomodulatory, muscular, vulnary, gastrointestinal,
 CC nephrotropic, antineoplastic, gynaecological and antibacterial activities,
 CC and can be used in gene therapy. The prostate cancer antigen
 CC polynucleotides may be used for detection of prostate cancer, chromosome
 CC identification, as chromosome markers, and for numerous other diagnostic
 CC or research purposes. The prostate cancer antigens may be used to treat
 CC disorders such as neural, immune, muscular, reproductive,
 CC gastrointestinal, pulmonary, cardiovascular, renal and proliferative
 CC disorders, wounds, and infectious diseases. AAF16506 to AAF16514 to
 CC AAB57303 represent sequences used in the exemplification of the present
 CC invention.
 XX
 SQ Sequence 88 AA;

Query Match 73.9%; Score 34; DB 21; Length 88;
 Best Local Similarity 62.5%; Pred. NO. 44;
 Matches 5; Conservative 3; Mismatches 0; Indels 0; Gaps 0;

QY 1 QKLCHQKK 8
 I:||||:
 DB 62 qeichqqe 69

RESULT 4
 AAG76544
 ID AAG76544 standard; Protein; 102 AA.

XX AAG76544;

XX 03-SEP-2001 (first entry)

DE Human colon cancer antigen protein SEQ ID NO:7308.

XX Human; colon cancer; colon cancer antigen; diagnosis; detection;
 KW colorectal carcinoma.

XX Homo sapiens.

XX WO200122920-A2.

XX 05-APR-2001.

XX 28-SEP-2000; 2000WO-US26524.

XX 29-SEP-1999; 99US-0157137.

XX 03-NOV-1999; 99US-0163280.

XX (HUMA-) HUMAN GENOME SCI INC.

PI Ruben SM, Barash SC, Birse CE, Rosen CA;

XX WPI: 2001-235357/24.

DR N-PSDB; AAB35949.

XX Nucleic acids encoding 4277 human colon cancer-associated polypeptides,
 PT useful for preventing, diagnosing and/or treating colorectal cancers -

PS Claim 11: Page 8741-8742; 9803pp; English.

XX AAB32943 to AAB37195 and AAG73514 to AAG77788 represent human colon
 CC cancer-associated nucleic acid molecules (N) and proteins (P), where
 CC the proteins are collectively known as colon cancer antigens. The colon
 CC cancer antigens have cytostatic activity and can be used in gene
 CC therapy and vaccine production. N and P may be used in the prevention,
 CC diagnosis and treatment of diseases associated with inappropriate P
 CC expression. For example, N and P may be used to treat disorders
 CC associated with decreased expression by rectifying mutations or deletions
 CC in a patient's genome that affect the activity of P by expressing
 CC inactive proteins or to supplement the patients own production of P.

CC Additionally, N may be used to produce the colon cancer-associated Ps,
 CC by inserting the nucleic acids into a host cell and culturing the cell
 CC to express the proteins. N and P can be used in the prevention, diagnosis
 CC and treatment of colorectal carcinomas and cancers. AAB37196 to AAB37204
 CC and AAB77789 represent sequences used in the exemplification of the
 CC present invention.
 CC N.B. Pages 666 to 682 and page 7053 of the sequence listing were
 CC missing at time of publication, meaning no sequences are present for
 CC SEQ ID NO:1027 to 1052, 7921 and 7922.

XX
 SQ Sequence 102 AA;

Query Match 73.9%; Score 34; DB 22; Length 102;
 Best Local Similarity 83.3%; Pred. NO. 50;
 Matches 5; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

QY 3 LCHQKK 8
 :|||||
 DB 61 lchqkk 66

RESULT 5

AAB95513

ID AAB95513 standard; Protein; 522 AA.

XX AAB95513;

XX 26-JUN-2001 (first entry)

DE Human protein sequence SEQ ID NO:18081.

XX Human; primer; detection; diagnosis; antisense therapy; gene therapy.

XX Homo sapiens.

XX EP1074617-A2.

XX 07-FEB-2001.

XX 28-JUL-2000; 2000EP-0116126.

XX 29-JUL-1999; 99JP-0248036.

XX 27-AUG-1999; 99JP-0300253.

XX 11-JAN-2000; 2000JP-0118776.

XX 02-MAY-2000; 2000JP-0183767.

XX 09-JUN-2000; 2000JP-0241899.

XX (HELI-) HELIX RES INST.

PI Ota T, Isogai T, Nishikawa T, Hayashi K, Saito K, Yamamoto J;
 PI Ishii S, Sugiyama T, Wakamatsu A, Nagai K, Otsuki T;

XX WPI: 2001-318749/34.

XX Primer sets for synthesizing polynucleotides, particularly the 5602
 PT full-length cDNAs defined in the specification, and for the detection
 PT and/or diagnosis of the abnormality of the proteins encoded by the
 PT full-length cDNAs -

XX Claim 8; SEQ ID 18081; 2537pp + CD ROM; English.

XX The present invention describes primer sets for synthesizing 5602
 CC full-length cDNAs defined in the specification. Where a primer set
 CC comprises: (a) an oligo-dT primer and an oligonucleotide complementary
 CC to the complementary strand of a polynucleotide which comprises one of
 CC the 5602 nucleotide sequences defined in the specification, where the
 CC oligonucleotide comprises at least 15 nucleotides; or (b) a combination
 CC of an oligonucleotide comprising a sequence complementary to the
 CC complementary strand of a polynucleotide which comprises a 5'-end
 CC sequence and an oligonucleotide comprising a sequence complementary to a
 CC polynucleotide which comprises a 3'-end sequence, where the
 CC oligonucleotide comprises at least 15 nucleotides and the combination of

CC the 5'-end sequence/3'-end sequence is selected from those defined in
 CC the specification. The primer sets can be used in antisense therapy and
 CC in gene therapy. The primers are useful for synthesizing polynucleotides,
 CC particularly full-length cDNAs. The primers are also useful for the
 CC detection and/or diagnosis of the abnormality of the proteins encoded by
 CC the full-length cDNAs. The primers allow obtaining of the full-length
 CC cDNAs easily without any specialised methods. AAH03166 to AAH13628 and
 CC AAH13633 to AAH18742 represent human cDNA sequences; AAB92446 to
 CC AAB95893 represent human amino acid sequences; and AAH13629 to AAH13632
 CC represent oligonucleotides, all of which are used in the exemplification
 CC of the present invention.

XX Sequence 522 AA;

Query Match 73.9%; Score 34; DB 22; Length 522;
 Best Local Similarity 62.5%; Pred. No. 2.2e+02;
 Matches 5; Conservative 3; Mismatches 0; Indels 0; Gaps 0;

QY 1 QKLCHQKK 8
 Db 157 qelchqge 164

RESULT 6

AAAB93045
 ID AAB93045 standard; Protein; 790 AA.

XX AAB93045;

XX 26-JUN-2001 (first entry)

DE Human protein sequence SEQ ID NO:11834.

XX Human; primer; detection; diagnosis; antisense therapy; gene therapy.

XX Homo sapiens.

XX EP1074617-A2.

XX 07-FEB-2001.

XX 28-JUL-2000; 2000EP-0116126.

XX 29-JUL-1999; 99JP-0248036.

XX 27-AUG-1999; 99JP-0300253.

XX 11-JAN-2000; 2000JP-0118776.

XX 02-MAY-2000; 2000JP-0183767.

XX 09-JUN-2000; 2000JP-0241899.

XX (HELI-) HELIX RES INST.

XX Ota T, Isegai T, Nishikawa T, Hayashi K, Saito K, Yamamoto J;
 PI Ishii S, Sugiyama T, Wakamatsu A, Nagai K, Otsuki T;

XX WPI; 2001-318749/34.

XX Primer sets for synthesizing polynucleotides, particularly the 5602
 PT full-length cDNAs defined in the specification, and for the detection
 PT and/or diagnosis of the abnormality of the proteins encoded by the
 PT full-length cDNAs -

XX Claim 8; SEQ ID 11834; 2537pp + CD ROM; English.

XX The present invention describes primer sets for synthesizing 5602
 CC full-length cDNAs defined in the specification. Where a primer set
 CC comprises: (a) an oligo-dT primer and an oligonucleotide complementary
 CC to the complementary strand of a polynucleotide which comprises one of
 CC the 5602 nucleotide sequences defined in the specification, where the
 CC oligonucleotide comprises at least 15 nucleotides; or (b) a combination
 CC of an oligonucleotide comprising a sequence complementary to the
 CC complementary strand of a polynucleotide which comprises a 5'-end
 CC sequence and an oligonucleotide comprising a sequence complementary to a

CC polynucleotide which comprises a 3'-end sequence, where the
 CC oligonucleotide comprises at least 15 nucleotides and the combination of
 CC the 5'-end sequence/3'-end sequence is selected from those defined in
 CC the specification. The primer sets can be used in antisense therapy and
 CC in gene therapy. The primers are useful for synthesizing polynucleotides,
 CC particularly full-length cDNAs. The primers are also useful for the
 CC detection and/or diagnosis of the abnormality of the proteins encoded by
 CC the full-length cDNAs. The primers allow obtaining of the full-length
 CC cDNAs easily without any specialised methods. AAH03166 to AAH13628 and
 CC AAH13633 to AAH18742 represent human cDNA sequences; AAB92446 to
 CC AAB95893 represent human amino acid sequences; and AAH13629 to AAH13632
 CC represent oligonucleotides, all of which are used in the exemplification
 CC of the present invention.

XX Sequence 790 AA;

Query Match 73.9%; Score 34; DB 22; Length 790;
 Best Local Similarity 62.5%; Pred. No. 3.2e+02;
 Matches 5; Conservative 3; Mismatches 0; Indels 0; Gaps 0;

QY 1 QKLCHQKK 8
 Db 157 qelchqge 164

RESULT 7

AAAB18183
 ID AAB18183 standard; Protein; 1188 AA.

XX AAB18183;

XX 07-NOV-2000 (first entry)

XX Plasmodium falciparum chromosome 2 related protein SEQ ID NO:40.

XX Plasmodium falciparum; chromosome 2; human malaria parasite; vaccine;
 XX antimalarial; malaria; protozoacide; infection; insecticide.

XX Plasmodium falciparum.

XX WO200025728-A2.

XX 11-MAY-2000.

XX 05-NOV-1999; 99WO-US26796.

XX 05-NOV-1998; 98US-0107131.

XX (HOFF/) HOFFMAN S.

XX (CARU/) CARUCCI D.

XX (GARD/) GARDNER M.

XX (VENT/) VENTER J C.

XX Hoffman S, Carucci D, Gardner M, Venter JC;

XX WPI; 2000-365347/31.

XX Proteins encoded by chromosome 2 of the human malarial parasite,
 PT Plasmodium falciparum, useful as antimalarial vaccines and in the
 PT diagnosis of P.falciparum infection -

XX Disclosure; Page 101-104; 577pp; English.

XX The present invention describes proteins and their fragments (I) encoded
 CC by chromosome 2 of the human malarial parasite, Plasmodium falciparum.
 CC Also described are: (1) nucleotide sequences (II) encoding (I); and (2)
 CC vaccines against P. falciparum infection comprising (I) or (II).
 CC (I) and (II) are useful for the development of vaccines against
 CC P. falciparum infection. (I) and polyclonal antisera or a monoclonal
 CC antibody raised to immunogens comprising the sequences of (I), are
 CC useful in the detection of infection with P. falciparum. Furthermore,
 CC (I) (especially when they are rifins or secreted or membrane proteins)

CC can aid the identification of drugs to treat or prevent P. falciparum
CC infection, or they can be used to identify drug resistance in
CC P. falciparum. Sequencing of the plasmodium chromosome 2 and the
CC subsequent identification of proteins encoded by it will help to expand
CC our understanding of parasite biology, a process hampered by the
CC complexity of the parasitic lifecycle, and provide new targets for
CC vaccine and drug development. Parasite resistance to drugs and mosquito
CC resistance to insecticides have led to a resurgence of malaria in many
CC parts of the world, and there is a pressing need for vaccines and new
CC drugs. AAA70078 to AAA70287 and AAB18144 to AAB18352 represent nucleotide
CC and protein sequences given in the present invention, but which are not
CC specifically mentioned within the specification.

XX Sequence 1188 AA;

Query Match 73.9%; Score 34; DB 21; Length 1188;
Best Local Similarity 71.4%; Pred. No. 4.6e+02;
Matches 5; Conservative 2; Mismatches 0; Indels 0; Gaps 0;

QY 1 QKICHQK 7

:|:||||

Db 1117 eelchqk 1123

RESULT 8

AAQ08675
ID AAG08675 standard; Protein; 197 AA.

XX AC AAG08675;

XX DT 17-OCT-2000 (first entry)

XX DE Arabidopsis thaliana protein fragment SEQ ID NO: 6307.

KW Protein identification; signal transduction pathway; metabolic pathway;
KW hybridisation assay; genetic mapping; gene expression control; promoter;
KW termination sequence.

XX OS Arabidopsis thaliana.

XX PN EPI033405-A2.

XX PD 06-SEP-2000.

XX PF 25-FEB-2000; 2000EP-0301439.

XX PR 25-FEB-1999; 99US-0121825.

XX PR 03-MAR-1999; 99US-0123180.

XX PR 09-MAR-1999; 99US-0123548.

XX PR 23-MAR-1999; 99US-0125788.

XX PR 25-MAR-1999; 99US-0126264.

XX PR 29-MAR-1999; 99US-0126785.

XX PR 01-APR-1999; 99US-0127462.

XX PR 06-APR-1999; 99US-0128234.

XX PR 16-APR-1999; 99US-0128714.

XX PR 16-APR-1999; 99US-0129845.

XX PR 19-APR-1999; 99US-0130077.

XX PR 21-APR-1999; 99US-0130449.

XX PR 23-APR-1999; 99US-0130510.

XX PR 28-APR-1999; 99US-0130891.

XX PR 30-APR-1999; 99US-0131449.

XX PR 30-APR-1999; 99US-0132048.

XX PR 04-MAY-1999; 99US-0132407.

XX PR 05-MAY-1999; 99US-0132484.

XX PR 05-MAY-1999; 99US-0132485.

XX PR 06-MAY-1999; 99US-0132486.

XX PR 06-MAY-1999; 99US-0132487.

XX PR 07-MAY-1999; 99US-0132863.

XX PR 11-MAY-1999; 99US-0134256.

XX PR 14-MAY-1999; 99US-0134218.

XX PR 14-MAY-1999; 99US-0134219.

XX PR 14-MAY-1999; 99US-0134221.

PR 14-MAY-1999; 99US-0134370.
PR 18-MAY-1999; 99US-0134768.
PR 19-MAY-1999; 99US-0134941.
PR 20-MAY-1999; 99US-0135124.
PR 21-MAY-1999; 99US-0135353.
PR 24-MAY-1999; 99US-0135629.
PR 25-MAY-1999; 99US-0136021.
PR 27-MAY-1999; 99US-0136392.
PR 28-MAY-1999; 99US-0136782.
PR 01-JUN-1999; 99US-0137222.
PR 03-JUN-1999; 99US-0137528.
PR 04-JUN-1999; 99US-0137502.
PR 07-JUN-1999; 99US-0137724.
PR 08-JUN-1999; 99US-0138094.
PR 10-JUN-1999; 99US-0138540.
PR 10-JUN-1999; 99US-0138847.
PR 14-JUN-1999; 99US-0139119.
PR 16-JUN-1999; 99US-0139452.
PR 16-JUN-1999; 99US-0139453.
PR 17-JUN-1999; 99US-0139492.
PR 18-JUN-1999; 99US-0139454.
PR 18-JUN-1999; 99US-0139455.
PR 18-JUN-1999; 99US-0139456.
PR 18-JUN-1999; 99US-0139457.
PR 18-JUN-1999; 99US-0139458.
PR 18-JUN-1999; 99US-0139459.
PR 18-JUN-1999; 99US-0139460.
PR 18-JUN-1999; 99US-0139461.
PR 18-JUN-1999; 99US-0139462.
PR 18-JUN-1999; 99US-0139463.
PR 18-JUN-1999; 99US-0139750.
PR 18-JUN-1999; 99US-0139763.
PR 21-JUN-1999; 99US-0139817.
PR 22-JUN-1999; 99US-0139899.
PR 23-JUN-1999; 99US-0140353.
PR 23-JUN-1999; 99US-0140354.
PR 24-JUN-1999; 99US-0140695.
PR 28-JUN-1999; 99US-0140823.
PR 29-JUN-1999; 99US-0140991.
PR 30-JUN-1999; 99US-0141287.
PR 01-JUL-1999; 99US-0141842.
PR 01-JUL-1999; 99US-0142154.
PR 02-JUL-1999; 99US-0142055.
PR 06-JUL-1999; 99US-0142390.
PR 08-JUL-1999; 99US-0142803.
PR 09-JUL-1999; 99US-0142920.
PR 12-JUL-1999; 99US-0142977.
PR 13-JUL-1999; 99US-0143542.
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PR 15-AUG-1999; 99US-0149368.
PR 17-AUG-1999; 99US-0149175.
PR 18-AUG-1999; 99US-0149426.
PR 20-AUG-1999; 99US-0149722.
PR 20-AUG-1999; 99US-0149723.
PR 23-AUG-1999; 99US-0149902.
PR 23-AUG-1999; 99US-0149930.
PR 25-AUG-1999; 99US-0150566.
PR 26-AUG-1999; 99US-0150884.
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PR 27-AUG-1999; 99US-0151066.
PR 27-AUG-1999; 99US-0151080.
PR 30-AUG-1999; 99US-0151303.
PR 31-AUG-1999; 99US-0151438.
PR 01-SEP-1999; 99US-0151930.
PR 07-SEP-1999; 99US-0152363.
PR 10-SEP-1999; 99US-0153070.
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PR 25-OCT-1999; 99US-0161406.
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PR 26-OCT-1999; 99US-0161360.
PR 26-OCT-1999; 99US-0161361.
PR 28-OCT-1999; 99US-0161920.
PR 28-OCT-1999; 99US-0161922.
PR 28-OCT-1999; 99US-0161993.
PR 28-OCT-1999; 99US-0162142.

Query Match 71.7%; Score 33; DB 21; Length 197;
Best Local Similarity 71.4%; Pred. No. 1.4e+02;
Matches 5; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

Qy 2 KLCHQKK 8
Db 185 klchkr 191

RESULT 9

AAG08674
ID AAG08674 standard; Protein; 242 AA.

XX
AC AAG08674;

XX
DT 17-OCT-2000 (first entry)

XX
DE Arabidopsis thaliana protein fragment SEQ ID NO: 6306.

KW Protein identification; signal transduction pathway; metabolic pathway;
KW hybridisation assay; genetic mapping; gene expression control; promoter;
KW termination sequence.

OS Arabidopsis thaliana.

XX
PN EF1033405-A2.

XX
PD 06-SEP-2000.

XX
PF 25-FEB-2000; 2000EP-0301439.

PR 25-FEB-1999; 99US-0121825.
PR 03-MAR-1999; 99US-0123180.
PR 09-MAR-1999; 99US-0123548.
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PR 25-MAY-1999; 99US-0136021.

PR 27-MAY-1999; 99US-0136392.
PR 28-MAY-1999; 99US-0136782.
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PR 26-AUG-1999; 99US-0150884.
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PR 30-AUG-1999; 99US-0151303.
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PR 06-OCT-1999; 99US-0157865.
PR 07-OCT-1999; 99US-0158029.
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PR 28-OCT-1999; 99US-0161992.
PR 28-OCT-1999; 99US-0161993.

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PR 29-OCT-1999; 99US-0162142.
Query Match 71.7%; Score 33; DB 21; Length 242;
Best Local Similarity 71.4%; Pred. No. 1.6e+02;
Matches 5; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

QY 2 KLCHQK 8
Db 230 klchkr 236
|||||:

RESULT 10
AAM25801
ID AAM25801 standard; Protein; 678 AA.
XX AC AAM25801;
XX DT 16-OCT-2001 (first entry)
XX DE Human protein sequence SPQ ID NO:1316.
XX KW Human; cancer; ulcer; HIV infection; human immunodeficiency virus;
KW antiinflammatory; antirheumatic; antiarthritic; immunosuppressive;
KW antibacterial; endocrine; cardiant; central nervous system; virucide;
KW anti-HIV; fungicide; antimutagen; cardiovascular; antianaemic; anaemia;
KW antiaggregant; haemostatic; vulnery; antilucer; osteopathic; eczema;
KW dermatological; anti-allergic; antiasthmatic; antidiabetic; cytostatic;
KW neuroprotective; antidepressant; nootropic; antiparkinsonian; infection;
KW immunostimulant; gene therapy; antisense therapy; vaccine; inflammation;
KW antianaphylactic; rheumatoid arthritis; septic shock; pancreatitis;
KW cardiac dysfunction; neuropathology; cardiac anaphylaxis; autoimmune;
KW genetic disease; haematopoietic disorder; platelet disorder; asthma;
KW thrombocytopaenia; osteoporosis; severe combined immunodeficiency;
KW allergic rhinitis; diabetes; multiple sclerosis; depression;
KW Alzheimer's disease; Parkinson's disease; neurodegenerative disorder;
KW neurological disorder.
XX OS Homo sapiens.
XX PN WO200153455-A2.
XX PD 26-JUL-2001.
XX PF 22-DEC-2000; 2000WO-US35017.
XX PR 23-DEC-1999; 99US-0471275.
XX PR 21-JAN-2000; 2000US-0488725.
XX PR 25-APR-2000; 2000US-052317.
XX PA (HYSE-) HYSEQ INC.
XX PI Tang YT, Liu C, Drmanac RT;
XX N-PSDB; AAH59742.
XX WPI: 2001-457603/49.
XX DR N-PSDB; AAH59742.
XX FT Isolated human polynucleotides encoding polypeptides, useful for the
XX treatment and diagnosis of e.g. cancer, ulcers and HIV infection.
XX PS Claim 20; Page 274; 1217pp; English.
XX CC AAH99166 to AAH99904 encode the human proteins given in AAM25225 to
XX AAM25963. The proteins can have activities based on the tissues and
XX cells they are expressed in, such as: antiinflammatory; antirheumatic;
XX antiarthritic; immunosuppressive; antibacterial; endocrine; cardiant;
XX central nervous system; virucide; anti-HIV; fungicide; antimutagen;
XX cardiovascular; antianaemic; antiaggregant; haemostatic; vulnery;
XX antilucer; osteopathic; dermatological; antiallergic; antiasthmatic;
XX antidiabetic; cytostatic; neuroprotective; antidepressant; nootropic;
XX antiparkinsonian; and immunostimulant. The proteins and polynucleotides
XX encoding them can be used in gene therapy, antisense therapy and vaccine
XX production. The proteins and polynucleotides are useful for screening for
XX agonists or antagonists of a protein and for the treatment and diagnosis
XX of disorders associated with the activity of a protein e.g. inflammation,
XX rheumatoid arthritis, septic shock, pancreatitis, viral, bacterial, HIV and fungal
XX neuropathology, cardiac anaphylaxis, genetic diseases, haematopoietic disorders,
XX infections, autoimmunity, genetic diseases, haematopoietic disorders,
XX anaemia, platelet disorders, thrombocytopaenia, wounds, burns, ulcers;
XX osteoporosis, severe combined immunodeficiency, eczema, allergic
XX rhinitis, asthma, diabetes, cancer, multiple sclerosis, depression,
XX Alzheimer's disease, Parkinson's disease, neurodegenerative and
XX neurological disorders.
XX SQ Sequence 678 AA;

Query Match 71.7%; Score 33; DB 22; Length 678;
Best Local Similarity 83.3%; Pred. No. 4.1e+02;
Matches 5; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

QY 2 KLCHQK 7
Db 610 klchkr 615
|||||:

RESULT 11
AAY79180
ID AAY79180 standard; Protein; 778 AA.
XX AC AAY79180;
XX DT 19-JUN-2000 (first entry)
XX DE Haematopoietic stem cell specific protein.
XX KW Haematopoietic stem cell; immune system disorder;
XX leukaemia; antileukaemic; immunomodulator; therapy; mouse.
XX OS Mus musculus.
XX FH Key Location/Qualifiers
XX FT Misc-difference 9 /note= "encoded by CAS"
XX FT Misc-difference 21 /note= "encoded by ASA"
XX FT Misc-difference 32 /note= "encoded by SAA"
XX FT Misc-difference 738 /note= "encoded by TTN"
XX PN WO200011168-A2.
XX PD 02-MAR-2000.
XX PF 20-AUG-1999; 99WO-US19052.
XX PR 21-AUG-1998; 98US-0138132.
XX PA (UYPR-) UNIV PRINCETON.
XX PI Lemischka I, Moore K;
XX WPI: 2000-237650/20.
XX DR N-PSDB; AAZ94121.
XX FT Hematopoietic stem cell signaling proteins modulating replication and
XX differentiation for treating immune system disorders and leukaemia.
XX PS Claim 21; Page 221-223; 256pp; English.
XX CC The present sequence is that of a mouse haematopoietic stem cell
XX (HSC) specific protein. It is an example of claimed HSC-specific
XX proteins (see AAY79176-93) predicted from novel isolated HSC-specific
XX nucleic acids (see AA294077-131). The HSCs are especially primitive
XX HSCs (PHSCs) such as umbilical cord cells, bone marrow cells and
XX foetal liver cells. The encoded proteins are growth factors,

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transcription factors, splicing factors, capping factors, transport proteins, translation factors or replication factors that modulate HSC activity, especially differentiation or replication. The invention provides claimed methods: for identifying PHSC-specific nucleic acids; for generating a stem cell/progenitor cell from PHSCs; for identifying the presence of a PHSC in a sample; for identifying the presence in a sample of a compound that modulates HSC activity; for using such a compound to treat an immune system condition, especially leukaemia; for introducing exogenous nucleic acid into a HSC; and for ex vivo expansion of HSCs. Also claimed are vectors, host cells, and an antibody that specifically binds a an HSC-specific protein.

XX SQ Sequence 778 AA;

Query Match 71.7%; Score 33; DB 21; Length 778;
Best Local Similarity 62.5%; Pred. No. 4.7e+02;
Matches 5; Conservative 1; Mismatches 2; Indels 0; Gaps 0;

QY 1 QKLCHQKK 8
| | | | |
Db 736 qsxchekk 743

RESULT 12
AAW79501
ID AAW79501 standard; peptide; 23 AA.
AC AAW79501;
XX
DT 17-DEC-1998 (first entry)
XX
DE Wild-type yeast transcription factor IIB (region 144-160).
XX
KW Yeast transcription factor IIB; YTFIIB; cell growth; mutation; hybrid;
KW human; antifungal drug.
XX
OS Saccharomyces cerevisiae.
XX
PN WO9839355-A1.
XX
PD 11-SEP-1998.
XX
PF 13-JUN-1997; 97WO-US10404.
XX
PR 06-MAR-1997; 97US-0812175.
XX
PA (CHIL-) CHILDREN'S HOSPITAL MEDICAL CENT.
XX
PI Carson DJ, Dorsey MJ, Ma J, Shaw S, Wingfield J;
XX
DR WPI; 1998-495788/42.
XX
PT Use of peptide fragments of yeast Transcription Factor IIB - to
PT identify compounds that inhibit fungal cell growth, especially to
PT screen for antifungal drugs active against Candida albicans
XX
PS Disclosure; Figure 5; 32pp; English.
XX
CC The present sequence represents the wild-type species-specific region of
CC the yeast transcription factor IIB (YTFIIB), amino acid residues 144-166.
CC This sequence has been found to contain 4 amino acids that are vital for
CC cell growth and in vivo activity of YTFIIB, mutations of these residues
CC results in cells having severe growth defects. By the analysis of hybrid
CC molecules in yeast cells, the 4 amino acids that confer yeast specificity
CC were identified as lysine 147, cysteine 149, lysine 151, and glutamic
CC acid 152. This wild-type sequence was used to generate loss-of-function
CC mutants, by changing a number of these critical amino acids to the
CC corresponding human residues. The YTFIIB fragment containing the vital 4
CC amino acid residues provides a yeast-specific target for screening
CC libraries to identify new antifungal drugs.

SQ Sequence 23 AA;

Query Match 69.6%; Score 32; DB 19; Length 23;
Best Local Similarity 71.4%; Pred. No. 29;
Matches 5; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

QY 2 KLCHQKK 8
| | | | |
Db 12 klchdek 18

RESULT 13
AAW79502
ID AAW79502 standard; peptide; 23 AA.
XX
AC AAW79502;
XX
DT 17-DEC-1998 (first entry)
XX
DE Loss-of-function mutant Yml.
XX
KW Yeast transcription factor IIB; YTFIIB; cell growth; mutation; hybrid;
KW human; antifungal drug.
XX
OS Saccharomyces cerevisiae.
OS Synthetic.
XX
FH Key Location/Qualifiers
FT Misc-difference 4 /note= "Wild-type Lys substituted by Val"
XX
PN WO9839355-A1.
XX
PD 11-SEP-1998.
XX
PF 13-JUN-1997; 97WO-US10404.
XX
PR 06-MAR-1997; 97US-0812175.
XX
PA (CHIL-) CHILDREN'S HOSPITAL MEDICAL CENT.
XX
PI Carson DJ, Dorsey MJ, Ma J, Shaw S, Wingfield J;
XX
DR WPI; 1998-495788/42.
XX
PT Use of peptide fragments of yeast Transcription Factor IIB - to
PT identify compounds that inhibit fungal cell growth, especially to
PT screen for antifungal drugs active against Candida albicans
XX
PS Disclosure; Figure 5; 32pp; English.
XX
CC Sequences AAW79502-W79510 are loss-of-function mutants, which were
CC generated by changing either one, three or all four of the vital 4 amino
CC acid residues of the YTFIIB species-specific wild-type sequence
CC (AAW79501) to the human equivalents. The four residues that confer
CC yeast specificity have been identified as lysine 147, cysteine 149,
CC lysine 151, and glutamic acid 152; these are vital for cell growth and in
CC vivo activity of YTFIIB, mutations of these residues results in cells
CC having severe growth defects. This particular YTFIIB derivative contains
CC a single mutation, however it was not seen to show any detectable
CC decrease in the protein's ability to support cell growth. The YTFIIB
CC fragment containing the vital 4 amino acid residues provides a
CC yeast-specific target for screening libraries to identify new antifungal
CC drugs.

XX SQ Sequence 23 AA;

Query Match 69.6%; Score 32; DB 19; Length 23;
Best Local Similarity 71.4%; Pred. No. 29;
Matches 5; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

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QY 2 KLCHOKK 8
DB 12 klchdex 18

RESULT 14
AAW79503
ID AAW79503 standard; peptide; 23 AA.
AC
XX
AC AAW79503;
DT 17-DEC-1998 (first entry)
XX
XX Loss-of-function mutant Ymc8.
DE
XX
XX Yeast transcription factor IIB; YTFIIB; cell growth; mutation; hybrid;
KW human; antifungal drug.
XX
XX Saccharomyces cerevisiae.
OS Synthetic.
XX
XX Key Location/Qualifiers
FH Misc-difference 6 /note= "Wild-type Cys substituted by Arg"
FT
XX
XX WO9839355-A1.
PN
XX 11-SEP-1998.
PD
XX 13-JUN-1997; 97WO-US10404.
PF
XX 06-MAR-1997; 97US-0812175.
PR
XX (CHIL-) CHILDREN'S HOSPITAL MEDICAL CENT.
PA
XX Carson DJ, Dorsey MJ, Ma J, Shaw S, Wingfield J;
PI WPI; 1998-495788/42.
DR
XX
XX Use of peptide fragments of yeast Transcription Factor IIB - to
PT identify compounds that inhibit fungal cell growth, especially to
PT screen for antifungal drugs active against Candida albicans
XX
PS Disclosure; Figure 5; 32pp; English.
XX
XX Sequences AAW79502-W79510 are loss-of-function mutants, which were
CC generated by changing either one, three or all four of the vital 4 amino
CC acid residues of the YTFIIB species-specific wild-type sequence
CC (AAW79501) to the human equivalents. The four residues that confer
CC yeast specificity have been identified as lysine 147, cysteine 149,
CC lysine 151, and glutamic acid 152, these are vital for cell growth and in
CC vivo activity of YTFIIB, mutations of these residues results in cells
CC having severe growth defects. This particular YTFIIB derivative contains
CC a single mutation, however it was not seen to show any detectable
CC decrease in the protein's ability to support cell growth. The YTFIIB
CC fragment containing the vital 4 amino acid residues provides a
CC yeast-specific target for screening libraries to identify new antifungal
CC drugs.
XX
SQ Sequence 23 AA;

Query Match 69.6%; Score 32; DB 19; Length 23;
Best Local Similarity 71.4%; Pred. No. 29;
Matches 5; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

QY 2 KLCHOKK 8
DB 12 klchdex 18

RESULT 15
AAW79504
ID AAW79504 standard; peptide; 23 AA.
AC
XX
AC AAW79504;
DT 17-DEC-1998 (first entry)
XX
XX Loss-of-function mutant Ymc8.
DE
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XX Yeast transcription factor IIB; YTFIIB; cell growth; mutation; hybrid;
KW human; antifungal drug.
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OS Synthetic.
XX
XX Key Location/Qualifiers
FH Misc-difference 8 /note= "Wild-type Lys substituted by Asn"
FT
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XX WO9839355-A1.
PN
XX 11-SEP-1998.
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DB 12 klchdex 18

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